

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 April 2001 (05.04.2001)

PCT

(10) International Publication Number
WO 01/23417 A2

- (51) International Patent Classification⁷: **C07K 14/00**
- (21) International Application Number: **PCT/EP00/09500**
- (22) International Filing Date:
27 September 2000 (27.09.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
9923154.0 30 September 1999 (30.09.1999) GB
0016839.3 7 July 2000 (07.07.2000) GB
- (71) Applicant (for all designated States except US):
SMITHKLINE BEECHAM BIOLOGICALS S.A.
[BE/BE]; Rue de l'Institut 89, B-1330 Rixensart (BE).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): **VINALS Y DE BAS-**
SOLS, Carlota [BE/BE]; SmithKline Beecham Biologi-
cals S.A., Rue de l'Institut 89, B-1330 Rixensart (BE).
- (74) Agent: **PRIVETT, Kathryn, Louise**; SmithKline
Beecham, Two New Horizons Court, Brentford, Middlesex
TW8 9EP (GB).
- (81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- Published:**
— Without international search report and to be republished
upon receipt of that report.
- For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.



WO 01/23417 A2

(54) Title: **NOVEL COMPOUNDS**

(57) Abstract: CASB6411 polypeptides and polynucleotides and methods for producing such polypeptides by recombinant techniques are disclosed. Also disclosed are methods for utilizing CASB6411 polypeptides and polynucleotides in diagnostics, and vaccines for prophylactic and therapeutic treatment of cancers, particularly ovarian and colon cancers, autoimmune diseases, and related conditions.

Novel Compounds

The present invention relates to polynucleotides, herein referred to as CASB6411 polynucleotides, polypeptides encoded thereby (referred to herein as CASB6411 polypeptides), recombinant materials and methods for their production. In another aspect, the invention relates to methods for using such polypeptides and polynucleotides, including the treatment of cancer and autoimmune diseases and other related conditions. In a further aspect, the invention relates to methods for identifying agonists and antagonists/inhibitors using the materials provided by the invention, and treating conditions associated with CASB6411 polypeptide imbalance with the identified compounds. In a still further aspect, the invention relates to diagnostic assays for detecting diseases associated with inappropriate CASB6411 polypeptide activity or levels.

Polypeptides and polynucleotides of the present invention are believed to be important immunogens for specific prophylactic or therapeutic immunization against tumours, because they are specifically expressed or highly over-expressed in tumours compared to normal cells and can thus be targeted by antigen-specific immune mechanisms leading to the destruction of the tumour cell. They can also be used to diagnose the occurrence of tumour cells. Furthermore, their inappropriate expression in certain circumstances can cause an induction of autoimmune, inappropriate immune responses, which could be corrected through appropriate vaccination using the same polypeptides or polynucleotides. In this respect the most important biological activities to our purpose are the antigenic and immunogenic activities of the polypeptide of the present invention. A polypeptide of the present invention may also exhibit at least one other biological activity of a CASB6411 polypeptide, which could qualify it as a target for therapeutic or prophylactic intervention different from that linked to the immune response.

In a first aspect, the present invention relates to CASB6411 polypeptides. Such peptides include isolated polypeptides comprising an amino acid sequence which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, most preferably at least 97-99% identity, to that of SEQ ID NO:2 OR 4 over the entire length of SEQ ID NO:2 OR 4. Such polypeptides include those comprising the amino acid of SEQ ID NO:2 OR 4.

Further peptides of the present invention include isolated polypeptides in which the amino acid sequence has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, most preferably at least 97-99% identity, to the amino acid sequence of SEQ ID NO:2 OR 4 over the entire length of SEQ ID NO:2 OR 4. Such polypeptides include the polypeptide of SEQ ID NO:2 OR 4.

Further peptides of the present invention include isolated polypeptides encoded by a polynucleotide comprising the sequence contained in SEQ ID NO:1.

The invention also provides an immunogenic fragment of a CASB6411 polypeptide, that is a contiguous portion of the CASB6411 polypeptide which has the same or similar immunogenic properties to the polypeptide comprising the amino acid sequence of SEQ ID NO:2 OR 4. That is to say, the fragment (if necessary when coupled to a carrier) is capable of raising an immune response which recognises the CASB6411 polypeptide. Such an immunogenic fragment may include, for example, the CASB6411 polypeptide lacking an N-terminal leader sequence, a transmembrane domain or a C-terminal anchor domain. In a preferred aspect the immunogenic fragment of CASB6411 according to the invention comprises substantially all of the extracellular domain of a polypeptide which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, most preferably at least 97-99% identity, to that of SEQ ID NO:2 OR 4 over the entire length of SEQ ID NO:2 OR 4. Preferably an immunogenic fragment according to the invention comprises at least one epitope.

Peptide fragments incorporating an epitope of CASB6411 typically will comprise at least 7, preferably 9 or 10 contiguous amino acids from SEQ ID NO:2 OR 4. Preferred epitopes are shown in SEQ ID NO:9 to SEQ ID NO:72.

Peptides that incorporate these epitopes form a preferred aspect of the present invention. Mimotopes which have the same characteristics as these epitopes, and immunogens comprising such mimotopes which generate an immune response which cross-react with

an epitope in the context of the CASB6411 molecule, also form part of the present invention.

The present invention, therefore, includes isolated peptides encompassing these epitopes themselves, and any mimotope thereof. The meaning of mimotope is defined as an entity
5 which is sufficiently similar to the native CASB6411 epitope so as to be capable of being recognised by antibodies which recognise the native molecule; (Gheysen, H.M., et al., 1986, Synthetic peptides as antigens. Wiley, Chichester, Ciba foundation symposium 119, p130-149; Gheysen, H.M., 1986, Molecular Immunology, 23,7, 709-715); or are
10 capable of raising antibodies, when coupled to a suitable carrier, which antibodies cross-react with the native molecule.

Peptide mimotopes of the above-identified epitopes may be designed for a particular purpose by addition, deletion or substitution of elected amino acids. Thus, the peptides of
15 the present invention may be modified for the purposes of ease of conjugation to a protein carrier. For example, it may be desirable for some chemical conjugation methods to include a terminal cysteine to the epitope. In addition it may be desirable for peptides conjugated to a protein carrier to include a hydrophobic terminus distal from the conjugated terminus of the peptide, such that the free unconjugated end of the peptide
20 remains associated with the surface of the carrier protein. This reduces the conformational degrees of freedom of the peptide, and thus increases the probability that the peptide is presented in a conformation which most closely resembles that of the peptide as found in the context of the whole molecule. For example, the peptides may be altered to have an N-terminal cysteine and a C-terminal hydrophobic amidated tail.

25 Alternatively, the addition or substitution of a D-stereoisomer form of one or more of the amino acids may be performed to create a beneficial derivative, for example to enhance stability of the peptide. Those skilled in the art will realise that such modified peptides, or mimotopes, could be a wholly or partly non-peptide mimotope wherein the constituent residues are not necessarily confined to the 20 naturally occurring amino acids. In
30 addition, these may be cyclised by techniques known in the art to constrain the peptide into a conformation that closely resembles its shape when the peptide sequence is in the context of the whole molecule. A preferred method of cyclising a peptide comprises the addition of a pair of cysteine residues to allow the formation of a disulphide bridge.

Further, those skilled in the art will realise that mimotopes or immunogens of the present invention may be larger than the above-identified epitopes, and as such may comprise the sequences disclosed herein. Accordingly, the mimotopes of the present invention may
5 consist of addition of N and/or C terminal extensions of a number of other natural residues at one or both ends. The peptide mimotopes may also be retro sequences of the natural sequences, in that the sequence orientation is reversed; or alternatively the sequences may be entirely or at least in part comprised of D-stereo isomer amino acids (inverso sequences). Also, the peptide sequences may be retro-inverso in character, in
10 that the sequence orientation is reversed and the amino acids are of the D-stereoisomer form. Such retro or retro-inverso peptides have the advantage of being non-self, and as such may overcome problems of self-tolerance in the immune system.

Alternatively, peptide mimotopes may be identified using antibodies which are capable
15 themselves of binding to the epitopes of the present invention using techniques such as phage display technology (EP 0 552 267 B1). This technique, generates a large number of peptide sequences which mimic the structure of the native peptides and are, therefore, capable of binding to anti-native peptide antibodies, but may not necessarily themselves share significant sequence homology to the native peptide. This approach may have
20 significant advantages by allowing the possibility of identifying a peptide with enhanced immunogenic properties, or may overcome any potential self-antigen tolerance problems which may be associated with the use of the native peptide sequence. Additionally this technique allows the identification of a recognition pattern for each native-peptide in terms of its shared chemical properties amongst recognised mimotope sequences.

25 The covalent coupling of the peptide to the immunogenic carrier can be carried out in a manner well known in the art. Thus, for example, for direct covalent coupling it is possible to utilise a carbodiimide, glutaraldehyde or (N-[γ -maleimidobutyryloxy] succinimide ester, utilising common commercially available heterobifunctional linkers
30 such as CDAP and SPDP (using manufacturers instructions). After the coupling reaction, the immunogen can easily be isolated and purified by means of a dialysis method, a gel filtration method, a fractionation method etc.

The types of carriers used in the immunogens of the present invention will be readily known to the man skilled in the art. The function of the carrier is to provide cytokine help in order to help induce an immune response against the peptide. A non-exhaustive list of carriers which may be used in the present invention include: Keyhole limpet

- 5 Haemocyanin (KLH), serum albumins such as bovine serum albumin (BSA), inactivated bacterial toxins such as tetanus or diphtheria toxins (TT and DT), or recombinant fragments thereof (for example, Domain 1 of Fragment C of TT, or the translocation domain of DT), or the purified protein derivative of tuberculin (PPD). Alternatively the mimotopes or epitopes may be directly conjugated to liposome carriers, which may
- 10 additionally comprise immunogens capable of providing T-cell help. Preferably the ratio of mimotopes to carrier is in the order of 1:1 to 20:1, and preferably each carrier should carry between 3-15 peptides.

- In an embodiment of the invention a preferred carrier is Protein D from *Haemophilus*
- 15 *influenzae* (EP 0 594 610 B1). Protein D is an IgD-binding protein from *Haemophilus influenzae* and has been patented by Forsgren (WO 91/18926, granted EP 0 594 610 B1). In some circumstances, for example in recombinant immunogen expression systems it may be desirable to use fragments of protein D, for example Protein D 1/3rd (comprising the N-terminal 100-110 amino acids of protein D (GB 9717953.5)).

20

Another preferred method of presenting the peptides of the present invention is in the context of a recombinant fusion molecule. For example, EP 0 421 635 B describes the use of chimaeric hepadnavirus core antigen particles to present foreign peptide sequences in a virus-like particle. As such, immunogens of the present invention may comprise peptides

25 presented in chimaeric particles consisting of hepatitis B core antigen. Additionally, the recombinant fusion proteins may comprise the mimotopes of the present invention and a carrier protein, such as NS1 of the influenza virus. For any recombinantly expressed protein which forms part of the present invention, the nucleic acid which encodes said immunogen also forms an aspect of the present invention.

30

Peptides used in the present invention can be readily synthesised by solid phase procedures well known in the art. Suitable syntheses may be performed by utilising "T-boc" or "F-moc" procedures. Cyclic peptides can be synthesised by the solid phase

procedure employing the well-known "F-moc" procedure and polyamide resin in the fully automated apparatus. Alternatively, those skilled in the art will know the necessary laboratory procedures to perform the process manually. Techniques and procedures for solid phase synthesis are described in 'Solid Phase Peptide Synthesis: A Practical
5 Approach' by E. Atherton and R.C. Sheppard, published by IRL at Oxford University Press (1989). Alternatively, the peptides may be produced by recombinant methods, including expressing nucleic acid molecules encoding the mimotopes in a bacterial or mammalian cell line, followed by purification of the expressed mimotope. Techniques for recombinant expression of peptides and proteins are known in the art, and are described
10 in Maniatis, T., Fritsch, E.F. and Sambrook et al., *Molecular cloning, a laboratory manual*, 2nd Ed.; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1989).

The polypeptides or immunogenic fragment of the invention may be in the form of the
15 "mature" protein or may be a part of a larger protein such as a precursor or a fusion protein. It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification such as multiple histidine residues, or an additional sequence for stability during recombinant production. Furthermore, addition of exogenous polypeptide or lipid
20 tail or polynucleotide sequences to increase the immunogenic potential of the final molecule is also considered.

In one aspect, the invention relates to genetically engineered soluble fusion proteins comprising a polypeptide of the present invention, or a fragment thereof, and various
25 portions of the constant regions of heavy or light chains of immunoglobulins of various subclasses (IgG, IgM, IgA, IgE). Preferred as an immunoglobulin is the constant part of the heavy chain of human IgG, particularly IgG1, where fusion takes place at the hinge region. In a particular embodiment, the Fc part can be removed simply by incorporation of a cleavage sequence which can be cleaved with blood clotting factor Xa. Furthermore,
30 this invention relates to processes for the preparation of these fusion proteins by genetic engineering, and to the use thereof for drug screening, diagnosis and therapy. A further aspect of the invention also relates to polynucleotides encoding such fusion proteins.

Examples of fusion protein technology can be found in International Patent Application Nos. WO94/29458 and WO94/22914.

The proteins may be chemically conjugated, or expressed as recombinant fusion proteins allowing increased levels to be produced in an expression system as compared to non-fused protein. The fusion partner may assist in providing T helper epitopes (immunological fusion partner), preferably T helper epitopes recognised by humans, or assist in expressing the protein (expression enhancer) at higher yields than the native recombinant protein. Preferably the fusion partner will be both an immunological fusion partner and expression enhancing partner.

Fusion partners include protein D from *Haemophilus influenza* B and the non-structural protein from influenzae virus, NS1 (hemagglutinin). Another immunological fusion partner is the protein known as LYTA. Preferably the C terminal portion of the molecule is used. Lyta is derived from *Streptococcus pneumoniae* which synthesize an N-acetyl-L-alanine amidase, amidase LYTA, (coded by the *lytA* gene {Gene, 43 (1986) page 265-272} an autolysin that specifically degrades certain bonds in the peptidoglycan backbone. The C-terminal domain of the LYTA protein is responsible for the affinity to the choline or to some choline analogues such as DEAE. This property has been exploited for the development of E.coli C-LYTA expressing plasmids useful for expression of fusion proteins. Purification of hybrid proteins containing the C-LYTA fragment at its amino terminus has been described {Biotechnology: 10, (1992) page 795-798}. It is possible to use the repeat portion of the Lyta molecule found in the C terminal end starting at residue 178, for example residues 188 - 305.

The present invention also includes variants of the aforementioned polypeptides, that is polypeptides that vary from the referents by conservative amino acid substitutions, whereby a residue is substituted by another with like characteristics. Typical such substitutions are among Ala, Val, Leu and Ile; among Ser and Thr; among the acidic residues Asp and Glu; among Asn and Gln; and among the basic residues Lys and Arg; or aromatic residues Phe and Tyr. Particularly preferred are variants in which several, 5-10, 1-5, 1-3, 1-2 or 1 amino acids are substituted, deleted, or added in any combination.

Polypeptides of the present invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood
5 in the art.

In a further aspect, the present invention relates to CASB6411 polynucleotides. Such polynucleotides include isolated polynucleotides comprising a nucleotide sequence encoding a polypeptide which has at least 70% identity, preferably at least 80% identity, more
10 preferably at least 90% identity, yet more preferably at least 95% identity, to the amino acid sequence of SEQ ID NO:2 OR 4, over the entire length of SEQ ID NO:2 OR 4. In this regard, polypeptides which have at least 97% identity are highly preferred, whilst those with at least 98-99% identity are more highly preferred, and those with at least 99% identity are most highly preferred. Such polynucleotides include a polynucleotide comprising the
15 nucleotide sequence contained in SEQ ID NO:1 or 3 encoding the polypeptide of SEQ ID NO:2 OR 4 respectively.

Further polynucleotides of the present invention include isolated polynucleotides comprising a nucleotide sequence that has at least 70% identity, preferably at least 80% identity, more
20 preferably at least 90% identity, yet more preferably at least 95% identity, to a nucleotide sequence encoding a polypeptide of SEQ ID NO:2 OR 4, over the entire coding region. In this regard, polynucleotides which have at least 97% identity are highly preferred, whilst those with at least 98-99% identity are more highly preferred, and those with at least 99% identity are most highly preferred.

25 Further polynucleotides of the present invention include isolated polynucleotides comprising a nucleotide sequence which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, to SEQ ID NO:1 or 3 or to the coding sequence of SEQ ID NO:1 or 3 over the entire
30 length of SEQ ID NO:1 or 3 or over the entire length of the coding sequence of SEQ ID NO:1 or 3 respectively. In this regard, polynucleotides which have at least 97% identity are highly preferred, whilst those with at least 98-99% identity are more highly preferred, and those with at least 99% identity are most highly preferred. Such polynucleotides include a

polynucleotide comprising the polynucleotide of SEQ ID NO:1 or 3 as well as the polynucleotide of SEQ ID NO:1 or 3 or the coding region of SEQ ID NO:1 or 3. Said polynucleotide can be inserted in a suitable plasmid or recombinant microorganism vector and used for immunization (see for example Wolff et. al., Science 247:1465-1468 (1990);
5 Corr et. al., J. Exp. Med. 184:1555-1560 (1996); Doe et. al., Proc. Natl. Acad. Sci. 93:8578-8583 (1996)).

The invention also provides polynucleotides which are complementary to all the above described polynucleotides.

10

The invention also provides a fragment of a CASB6411 polynucleotide which when administered to a subject has the same immunogenic properties as the polynucleotide of SEQ ID NO:1 or 3.

15 The invention also provides a polynucleotide encoding an immunological fragment of a CASB6411 polypeptide as hereinbefore defined.

The fragments have a level of immunogenic activity of at least about 50%, preferably at least about 70% and more preferably at least about 90% of the level of immunogenic
20 activity of a polypeptide sequence set forth in SEQ ID NO:2 OR 4 or a polypeptide sequence encoded by a polynucleotide sequence set forth in SEQ ID NO: 1.

The polypeptide fragments according to the invention preferably comprise at least about 5, 10, 15, 20, 25, 50, or 100 contiguous amino acids, or more, including all intermediate
25 lengths, of a polypeptide composition set forth herein, such as those set forth in SEQ ID NO:2 OR 4, or those encoded by a polynucleotide sequence set forth in a sequence of SEQ ID NO:1.

The nucleotide sequence of SEQ ID NO:1 is a cDNA sequence which comprises a
30 polypeptide encoding sequence (nucleotide 349 to 1761) encoding a polypeptide of 460 amino acids, the polypeptide of SEQ ID NO:2. The nucleotide sequence encoding the polypeptide of SEQ ID NO:2 may be identical to the polypeptide encoding sequence contained in SEQ ID NO:1 or it may be a sequence other than the one contained in SEQ

ID NO:1, which, as a result of the redundancy (degeneracy) of the genetic code, also encodes the polypeptide of SEQ ID NO:2. The polypeptide of the SEQ ID NO:2 is structurally related to other proteins of the LAK-4p family, having homology and/or structural similarity with Homo sapiens LAK-4p (GenBank accession BAA24179).

5

SEQ ID NO:3 is a cDNA sequence which is an alternative isoform of CASB6411 SEQ ID NO:1, and probably the result of alternative exon splicing. SEQ ID NO:3 comprises a polypeptide encoding sequence (nucleotide 382 to 844) encoding a polypeptide of 154 aminoacids, the polypeptide of SEQ ID NO:4. SEQ ID NO:4 is a truncated form of SEQ ID NO:2 polypeptide. The nucleotide sequence encoding the polypeptide of SEQ ID NO:4 may be identical to the polypeptide encoding sequence contained in SEQ ID NO:3 or it may be a sequence other than the one contained in SEQ ID NO:3, which, as a result of the redundancy (degeneracy) of the genetic code, also encodes the polypeptide of SEQ ID NO:4.

15

Preferred polypeptides and polynucleotides of the present invention are expected to have, *inter alia*, similar biological functions/properties to their homologous polypeptides and polynucleotides. Furthermore, preferred polypeptides, immunological fragments and polynucleotides of the present invention have at least one activity of either SEQ ID NO:1 or 3 or SEQ ID NO:2 OR 4, as appropriate.

20

The present invention also relates to partial or other incomplete polynucleotide and polypeptide sequences which were first identified prior to the determination of the corresponding full length sequences of SEQ ID NO:1 or 3 and SEQ ID NO:2 OR 4.

25

Accordingly, in a further aspect, the present invention provides for an isolated polynucleotide which:

(a) comprises a nucleotide sequence which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, even more preferably at least 97-99% identity to SEQ ID NO:5 and 7 over the entire length of SEQ ID NO:5 and 7, respectively;

30

(b) has a nucleotide sequence which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity,

even more preferably at least 97-99% identity, to SEQ ID NO:1 or 3 over the entire length of SEQ ID NO:5 and 7;

(c) the polynucleotide of SEQ ID NO:5 and 7; or

- (d) a nucleotide sequence encoding a polypeptide which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, even more preferably at least 97-99% identity, to the amino acid sequence of SEQ ID NO:6 and 8, over the entire length of SEQ ID NO:6 and 8, respectively,;
as well as the polynucleotide of SEQ ID NO:5 and 7.

10

The present invention further provides for a polypeptide which:

- (a) comprises an amino acid sequence which has at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, most preferably at least 97-99% identity, to that of SEQ ID NO:6 OR 8 over the entire length of SEQ ID NO:6 and 8;

- (b) has an amino acid sequence which is at least 70% identity, preferably at least 80% identity, more preferably at least 90% identity, yet more preferably at least 95% identity, most preferably at least 97-99% identity, to the amino acid sequence of SEQ ID NO:6 OR 8 over the entire length of SEQ ID NO:6 and 8;

- (c) comprises the amino acid of SEQ ID NO:6 and 8; and

(d) is the polypeptide of SEQ ID NO:6 and 8;

as well as polypeptides encoded by a polynucleotide comprising the sequence contained in SEQ ID NO:5 and 7.

- Polynucleotides of the present invention may be obtained, using standard cloning and screening techniques, from a cDNA library derived from mRNA in cells of human, colorectal tumours, stomach tumors and normal stomach, normal pancreas and pancreas tumors, ovarian tumors, lung tumors, and normal prostate, (for example Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Ed., Cold Spring harbor Laboratory Press, Cold Spring harbor, N.Y. (1989)). Polynucleotides of the invention can also be obtained from natural sources such as genomic DNA libraries or can be synthesized using well known and commercially available techniques.

When polynucleotides of the present invention are used for the recombinant production of polypeptides of the present invention, the polynucleotide may include the coding sequence for the mature polypeptide, by itself; or the coding sequence for the mature polypeptide in reading frame with other coding sequences, such as those encoding a leader or secretory
5 sequence, a pre-, or pro- or prepro- protein sequence, or other fusion peptide portions. For example, a marker sequence which facilitates purification of the fused polypeptide can be encoded. In certain preferred embodiments of this aspect of the invention, the marker sequence is a hexa-histidine peptide, as provided in the pQE vector (Qiagen, Inc.) and described in Gentz *et al.*, Proc Natl Acad Sci USA (1989) 86:821-824, or is an HA tag. The
10 polynucleotide may also contain non-coding 5' and 3' sequences, such as transcribed, non-translated sequences, splicing and polyadenylation signals, ribosome binding sites and sequences that stabilize mRNA.

Further embodiments of the present invention include polynucleotides encoding polypeptide
15 variants which comprise the amino acid sequence of SEQ ID NO:2 OR 4 and in which several, for instance from 5 to 10, 1 to 5, 1 to 3, 1 to 2 or 1, amino acid residues are substituted, deleted or added, in any combination.

Polynucleotides which are identical or sufficiently identical to a nucleotide sequence
20 contained in SEQ ID NO:1 or 3, may be used as hybridization probes for cDNA and genomic DNA or as primers for a nucleic acid amplification (PCR) reaction, to isolate full-length cDNAs and genomic clones encoding polypeptides of the present invention and to isolate cDNA and genomic clones of other genes (including genes encoding paralogs from human sources and orthologs and paralogs from species other than human) that have a high
25 sequence similarity to SEQ ID NO:1 or 3. Typically these nucleotide sequences are 70% identical, preferably 80% identical, more preferably 90% identical, most preferably 95% identical to that of the referent. The probes or primers will generally comprise at least 15 nucleotides, preferably, at least 30 nucleotides and may have at least 50 nucleotides. Particularly preferred probes will have between 30 and 50 nucleotides. Particularly
30 preferred primers will have between 20 and 25 nucleotides. In particular, polypeptides or polynucleotides derived from sequences from homologous animal origin could be used as immunogens to obtain a cross-reactive immune response to the human gene.

A polynucleotide encoding a polypeptide of the present invention, including homologs from species other than human, may be obtained by a process which comprises the steps of screening an appropriate library under stringent hybridization conditions with a labeled probe having the sequence of SEQ ID NO: 1 or 3 or a fragment thereof; and isolating full-length cDNA and genomic clones containing said polynucleotide sequence. Such hybridization techniques are well known to the skilled artisan. Preferred stringent hybridization conditions include overnight incubation at 42°C in a solution comprising: 50% formamide, 5xSSC (150mM NaCl, 15mM trisodium citrate), 50 mM sodium phosphate (pH7.6), 5x Denhardt's solution, 10 % dextran sulfate, and 20 microgram/ml denatured, sheared salmon sperm DNA; followed by washing the filters in 0.1x SSC at about 65°C. Thus the present invention also includes polynucleotides obtainable by screening an appropriate library under stringent hybridization conditions with a labeled probe having the sequence of SEQ ID NO:1 or 3 or a fragment thereof.

The skilled artisan will appreciate that, in many cases, an isolated cDNA sequence will be incomplete, in that the region coding for the polypeptide is short at the 5' end of the cDNA.

There are several methods available and well known to those skilled in the art to obtain full-length cDNAs, or extend short cDNAs, for example those based on the method of Rapid Amplification of cDNA ends (RACE) (see, for example, Frohman et al., PNAS USA 85, 8998-9002, 1988). Recent modifications of the technique, exemplified by the Marathon™ technology (Clontech Laboratories Inc.) for example, have significantly simplified the search for longer cDNAs. In the Marathon™ technology, cDNAs have been prepared from mRNA extracted from a chosen tissue and an 'adaptor' sequence ligated onto each end. Nucleic acid amplification (PCR) is then carried out to amplify the 'missing' 5' end of the cDNA using a combination of gene specific and adaptor specific oligonucleotide primers. The PCR reaction is then repeated using 'nested' primers, that is, primers designed to anneal within the amplified product (typically an adaptor specific primer that anneals further 3' in the adaptor sequence and a gene specific primer that anneals further 5' in the known gene sequence). The products of this reaction can then be analysed by DNA sequencing and a full-length cDNA constructed either by joining the

product directly to the existing cDNA to give a complete sequence, or carrying out a separate full-length PCR using the new sequence information for the design of the 5' primer.

- 5 Recombinant polypeptides of the present invention may be prepared by processes well known in the art from genetically engineered host cells comprising expression systems. Accordingly, in a further aspect, the present invention relates to an expression system which comprises a polynucleotide of the present invention, to host cells which are genetically engineered with such expression systems and to the production of polypeptides of the invention by recombinant techniques. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention.

- For recombinant production, host cells can be genetically engineered to incorporate expression systems or portions thereof for polynucleotides of the present invention. Introduction of polynucleotides into host cells can be effected by methods described in many standard laboratory manuals, such as Davis et al., Basic Methods in Molecular Biology (1986) and Sambrook *et al.*, Molecular Cloning: A Laboratory Manual, 2nd Ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (1989). Preferred such methods include, for instance, calcium phosphate transfection, DEAE-dextran mediated transfection, transfection, microinjection, cationic lipid-mediated transfection, electroporation, transduction, scrape loading, ballistic introduction or infection.

- Preferably the proteins of the invention are coexpressed with thioredoxin in trans (TIT). Coexpression of thioredoxin in trans versus in cis is preferred to keep antigen free of thioredoxin without the need for protease. Thioredoxin coexpression eases the solubilisation of the proteins of the invention. Thioredoxin coexpression has also a significant impact on protein purification yield, on purified-protein solubility and quality.

- 30 Representative examples of appropriate hosts include bacterial cells, such as *Streptococci*, *Staphylococci*, *E. coli*, *Streptomyces* and *Bacillus subtilis* cells; fungal cells, such as yeast cells and *Aspergillus* cells; insect cells such as *Drosophila* S2 and *Spodoptera* Sf9 cells;

animal cells such as CHO, COS, HeLa, C127, 3T3, BHK, HEK 293 and Bowes melanoma cells; and plant cells.

A great variety of expression systems can be used, for instance, chromosomal, episomal
5 and virus-derived systems, e.g., vectors derived from bacterial plasmids, from
bacteriophage, from transposons, from yeast episomes, from insertion elements, from
yeast chromosomal elements, from viruses such as baculoviruses, papova viruses, such as
SV40, vaccinia viruses, adenoviruses, fowl pox viruses, pseudorabies viruses and
retroviruses, and vectors derived from combinations thereof, such as those derived from
10 plasmid and bacteriophage genetic elements, such as cosmids and phagemids. The
expression systems may contain control regions that regulate as well as engender
expression. Generally, any system or vector which is able to maintain, propagate or
express a polynucleotide to produce a polypeptide in a host may be used. The appropriate
nucleotide sequence may be inserted into an expression system by any of a variety of
15 well-known and routine techniques, such as, for example, those set forth in Sambrook *et*
al., Molecular Cloning, A Laboratory Manual (supra). Appropriate secretion signals may
be incorporated into the desired polypeptide to allow secretion of the translated protein
into the lumen of the endoplasmic reticulum, the periplasmic space or the extracellular
environment. These signals may be endogenous to the polypeptide or they may be
20 heterologous signals.

The expression system may also be a recombinant live microorganism, such as a virus or
bacterium. The gene of interest can be inserted into the genome of a live recombinant
virus or bacterium. Inoculation and *in vivo* infection with this live vector will lead to *in*
25 *vivo* expression of the antigen and induction of immune responses.

Therefore, in certain embodiments, polynucleotides encoding immunogenic polypeptides
of the present invention are introduced into suitable mammalian host cells for expression
using any of a number of known viral-based systems. In one illustrative embodiment,
30 retroviruses provide a convenient and effective platform for gene delivery systems. A
selected nucleotide sequence encoding a polypeptide of the present invention can be
inserted into a vector and packaged in retroviral particles using techniques known in the
art. The recombinant virus can then be isolated and delivered to a subject. A number of

illustrative retroviral systems have been described (e.g., U.S. Pat. No. 5,219,740; Miller and Rosman (1989) *BioTechniques* 7:980-990; Miller, A. D. (1990) *Human Gene Therapy* 1:5-14; Scarpa et al. (1991) *Virology* 180:849-852; Burns et al. (1993) *Proc. Natl. Acad. Sci. USA* 90:8033-8037; and Boris-Lawrie and Temin (1993) *Cur. Opin. Genet. Develop.* 3:102-109.

In addition, a number of illustrative adenovirus-based systems have also been described. Unlike retroviruses which integrate into the host genome, adenoviruses persist extrachromosomally thus minimizing the risks associated with insertional mutagenesis (Haj-Ahmad and Graham (1986) *J. Virol.* 57:267-274; Bett et al. (1993) *J. Virol.* 67:5911-5921; Mittereder et al. (1994) *Human Gene Therapy* 5:717-729; Seth et al. (1994) *J. Virol.* 68:933-940; Barr et al. (1994) *Gene Therapy* 1:51-58; Berkner, K. L. (1988) *BioTechniques* 6:616-629; and Rich et al. (1993) *Human Gene Therapy* 4:461-476).

Various adeno-associated virus (AAV) vector systems have also been developed for polynucleotide delivery. AAV vectors can be readily constructed using techniques well known in the art. See, e.g., U.S. Pat. Nos. 5,173,414 and 5,139,941; International Publication Nos. WO 92/01070 and WO 93/03769; Lebkowski et al. (1988) *Molec. Cell. Biol.* 8:3988-3996; Vincent et al. (1990) *Vaccines* 90 (Cold Spring Harbor Laboratory Press); Carter, B. J. (1992) *Current Opinion in Biotechnology* 3:533-539; Muzyczka, N. (1992) *Current Topics in Microbiol. and Immunol.* 158:97-129; Kotin, R. M. (1994) *Human Gene Therapy* 5:793-801; Shelling and Smith (1994) *Gene Therapy* 1:165-169; and Zhou et al. (1994) *J. Exp. Med.* 179:1867-1875.

Additional viral vectors useful for delivering the nucleic acid molecules encoding polypeptides of the present invention by gene transfer include those derived from the pox family of viruses, such as vaccinia virus and avian poxvirus. By way of example, vaccinia virus recombinants expressing the novel molecules can be constructed as follows. The DNA encoding a polypeptide is first inserted into an appropriate vector so that it is adjacent to a vaccinia promoter and flanking vaccinia DNA sequences, such as the sequence encoding thymidine kinase (TK). This vector is then used to transfect cells

which are simultaneously infected with vaccinia. Homologous recombination serves to insert the vaccinia promoter plus the gene encoding the polypeptide of interest into the viral genome. The resulting TK.sup.(-) recombinant can be selected by culturing the cells in the presence of 5-bromodeoxyuridine and picking viral plaques resistant thereto.

5

- A vaccinia-based infection/transfection system can be conveniently used to provide for inducible, transient expression or coexpression of one or more polypeptides described herein in host cells of an organism. In this particular system, cells are first infected in vitro with a vaccinia virus recombinant that encodes the bacteriophage T7 RNA
- 10 polymerase. This polymerase displays exquisite specificity in that it only transcribes templates bearing T7 promoters. Following infection, cells are transfected with the polynucleotide or polynucleotides of interest, driven by a T7 promoter. The polymerase expressed in the cytoplasm from the vaccinia virus recombinant transcribes the transfected DNA into RNA which is then translated into polypeptide by the host
- 15 translational machinery. The method provides for high level, transient, cytoplasmic production of large quantities of RNA and its translation products. See, e.g., Elroy-Stein and Moss, Proc. Natl. Acad. Sci. USA (1990) 87:6743-6747; Fuerst et al. Proc. Natl. Acad. Sci. USA (1986) 83:8122-8126.
- 20 Alternatively, avipoxviruses, such as the fowlpox and canarypox viruses, can also be used to deliver the coding sequences of interest. Recombinant avipox viruses, expressing immunogens from mammalian pathogens, are known to confer protective immunity when administered to non-avian species. The use of an Avipox vector is particularly desirable in human and other mammalian species since members of the Avipox genus can only
- 25 productively replicate in susceptible avian species and therefore are not infective in mammalian cells. Methods for producing recombinant Avipoxviruses are known in the art and employ genetic recombination, as described above with respect to the production of vaccinia viruses. See, e.g., WO 91/12882; WO 89/03429; and WO 92/03545.
- 30 Any of a number of alphavirus vectors can also be used for delivery of polynucleotide compositions of the present invention, such as those vectors described in U.S. Patent Nos.

5,843,723; 6,015,686; 6,008,035 and 6,015,694. Certain vectors based on Venezuelan Equine Encephalitis (VEE) can also be used, illustrative examples of which can be found in U.S. Patent Nos. 5,505,947 and 5,643,576.

- 5 Moreover, molecular conjugate vectors, such as the adenovirus chimeric vectors described in Michael et al. *J. Biol. Chem.* (1993) 268:6866-6869 and Wagner et al. *Proc. Natl. Acad. Sci. USA* (1992) 89:6099-6103, can also be used for gene delivery under the invention.
- 10 Additional illustrative information on these and other known viral-based delivery systems can be found, for example, in Fisher-Hoch et al., *Proc. Natl. Acad. Sci. USA* 86:317-321, 1989; Flexner et al., *Ann. N.Y. Acad. Sci.* 569:86-103, 1989; Flexner et al., *Vaccine* 8:17-21, 1990; U.S. Patent Nos. 4,603,112, 4,769,330, and 5,017,487; WO 89/01973; U.S. Patent No. 4,777,127; GB 2,200,651; EP 0,345,242; WO 91/02805; Berkner,
- 15 *Biotechniques* 6:616-627, 1988; Rosenfeld et al., *Science* 252:431-434, 1991; Kolls et al., *Proc. Natl. Acad. Sci. USA* 91:215-219, 1994; Kass-Eisler et al., *Proc. Natl. Acad. Sci. USA* 90:11498-11502, 1993; Guzman et al., *Circulation* 88:2838-2848, 1993; and Guzman et al., *Cir. Res.* 73:1202-1207, 1993.
- 20 The recombinant live microorganisms described above can be virulent, or attenuated in various ways in order to obtain live vaccines. Such live vaccines also form part of the invention.
- 25 In certain embodiments, a polynucleotide may be integrated into the genome of a target cell. This integration may be in the specific location and orientation *via* homologous recombination (gene replacement) or it may be integrated in a random, non-specific location (gene augmentation). In yet further embodiments, the polynucleotide may be stably maintained in the cell as a separate, episomal segment of DNA. Such
- 30 polynucleotide segments or "episomes" encode sequences sufficient to permit maintenance and replication independent of or in synchronization with the host cell cycle.

The manner in which the expression construct is delivered to a cell and where in the cell the polynucleotide remains is dependent on the type of expression construct employed.

5 In another embodiment of the invention, a polynucleotide is administered/delivered as "naked" DNA, for example as described in Ulmer et al., *Science* 259:1745-1749, 1993 and reviewed by Cohen, *Science* 259:1691-1692, 1993. The uptake of naked DNA may be increased by coating the DNA onto biodegradable beads, which are efficiently transported into the cells.

10 In still another embodiment, a composition of the present invention can be delivered via a particle bombardment approach, many of which have been described. In one illustrative example, gas-driven particle acceleration can be achieved with devices such as those manufactured by Powderject Pharmaceuticals PLC (Oxford, UK) and Powderject Vaccines Inc. (Madison, WI), some examples of which are described in U.S. Patent Nos.
15 5,846,796; 6,010,478; 5,865,796; 5,584,807; and EP Patent No. 0500 799. This approach offers a needle-free delivery approach wherein a dry powder formulation of microscopic particles, such as polynucleotide or polypeptide particles, are accelerated to high speed within a helium gas jet generated by a hand held device, propelling the particles into a target tissue of interest.

20

In a related embodiment, other devices and methods that may be useful for gas-driven needle-less injection of compositions of the present invention include those provided by Bioject, Inc. (Portland, OR), some examples of which are described in U.S. Patent Nos. 4,790,824; 5,064,413; 5,312,335; 5,383,851; 5,399,163; 5,520,639 and 5,993,412.

25

Polypeptides of the present invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite
30 chromatography and lectin chromatography. Most preferably, ion metal affinity chromatography (IMAC) is employed for purification. Well known techniques for

refolding proteins may be employed to regenerate active conformation when the polypeptide is denatured during intracellular synthesis, isolation and or purification.

Another important aspect of the invention relates to a method for inducing , re-inforcing
5 or modulating an immunological response in a mammal which comprises inoculating the mammal with a fragment or the entire polypeptide or polynucleotide of the invention, adequate to produce antibody and/or T cell immune response for prophylaxis or for therapeutic treatment of cancer and autoimmune disease and related conditions. Yet another aspect of the invention relates to a method of inducing, re-inforcing or
10 modulating immunological response in a mammal which comprises, delivering a polypeptide of the present invention *via* a vector or cell directing expression of the polynucleotide and coding for the polypeptide *in vivo* in order to induce such an immunological response to produce immune responses for prophylaxis or treatment of said mammal from diseases.

15 A further aspect of the invention relates to an immunological/vaccine formulation (composition) which, when introduced into a mammalian host, induces, re-inforces or modulates an immunological response in that mammal to a polypeptide of the present invention wherein the composition comprises a polypeptide or polynucleotide of the
20 invention or an immunological fragment thereof as herein before defined. The vaccine formulation may further comprise a suitable carrier. Since a polypeptide may be broken down in the stomach, it is preferably administered parenterally (for instance, subcutaneous, intramuscular, intravenous, or intradermal injection). Formulations suitable for parenteral administration include aqueous and non-aqueous sterile injection
25 solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents or thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example, sealed ampoules and vials and may be stored in a freeze-dried condition requiring only the
30 addition of the sterile liquid carrier immediately prior to use.

A further aspect of the invention relates to the *in vitro* induction of immune responses to a fragment or the entire polypeptide or polynucleotide of the present invention or a

molecule comprising the polypeptide or polynucleotide of the present invention, using cells from the immune system of a mammal, and reinfusing these activated immune cells of the mammal for the treatment of disease. Activation of the cells from the immune system is achieved by *in vitro* incubation with the entire polypeptide or polynucleotide of the present invention or a molecule comprising the polypeptide or polynucleotide of the present invention in the presence or absence of various immunomodulator molecules.

A further aspect of the invention relates to the immunization of a mammal by administration of antigen presenting cells modified by *in vitro* loading with part or the entire polypeptide of the present invention or a molecule comprising the polypeptide of the present invention and administered *in vivo* in an immunogenic way. Alternatively, antigen presenting cells can be transfected *in vitro* with a vector containing a fragment or the entire polynucleotide of the present invention or a molecule comprising the polynucleotide of the present invention, such as to express the corresponding polypeptide, and administered *in vivo* in an immunogenic way.

15

According to another embodiment, the pharmaceutical compositions described herein will comprise one or more immunostimulants in addition to the immunogenic polynucleotide, polypeptide, antibody, T-cell and/or antigen presenting cell (APC) compositions of this invention. An immunostimulant refers to essentially any substance that enhances or
5 potentiates an immune response (antibody and/or cell-mediated) to an exogenous antigen. One preferred type of immunostimulant comprises an adjuvant. Many adjuvants contain a substance designed to protect the antigen from rapid catabolism, such as aluminum hydroxide or mineral oil, and a stimulator of immune responses, such as lipid A, *Bordetella pertussis* or *Mycobacterium tuberculosis* derived proteins. Certain adjuvants
10 are commercially available as, for example, Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, MI); Merck Adjuvant 65 (Merck and Company, Inc., Rahway, NJ); AS-2 (SmithKline Beecham, Philadelphia, PA); aluminum salts such as aluminum hydroxide gel (alum) or aluminum phosphate; salts of calcium, iron or zinc; an insoluble suspension of acylated tyrosine; acylated sugars; cationically or anionically
15 derivatized polysaccharides; polyphosphazenes; biodegradable microspheres; monophosphoryl lipid A and quil A. Cytokines, such as GM-CSF, interleukin-2, -7, -12, and other like growth factors, may also be used as adjuvants.

Within certain embodiments of the invention, the adjuvant composition is preferably one
20 that induces an immune response predominantly of the Th1 type. High levels of Th1-type cytokines (e.g., IFN- γ , TNF α , IL-2 and IL-12) tend to favor the induction of cell mediated immune responses to an administered antigen. In contrast, high levels of Th2-type cytokines (e.g., IL-4, IL-5, IL-6 and IL-10) tend to favor the induction of humoral immune responses. Following application of a vaccine as provided herein, a patient will
25 support an immune response that includes Th1- and Th2-type responses. Within a preferred embodiment, in which a response is predominantly Th1-type, the level of Th1-type cytokines will increase to a greater extent than the level of Th2-type cytokines. The levels of these cytokines may be readily assessed using standard assays. For a review of the families of cytokines, see Mosmann and Coffman, *Ann. Rev. Immunol.* 7:145-173,
30 1989.

Certain preferred adjuvants for eliciting a predominantly Th1-type response include, for example, a combination of monophosphoryl lipid A, preferably 3-de-O-acylated monophosphoryl lipid A, together with an aluminum salt. MPL[®] adjuvants are available from Corixa Corporation (Seattle, WA; *see*, for example, US Patent Nos. 4,436,727; 4,877,611; 4,866,034 and 4,912,094). CpG-containing oligonucleotides (in which the CpG dinucleotide is unmethylated) also induce a predominantly Th1 response. Such oligonucleotides are well known and are described, for example, in WO 96/02555, WO 99/33488 and U.S. Patent Nos. 6,008,200 and 5,856,462. Immunostimulatory DNA sequences are also described, for example, by Sato et al., *Science* 273:352, 1996.

Another preferred adjuvant comprises a saponin, such as Quil A, or derivatives thereof, including QS21 and QS7 (Aquila Biopharmaceuticals Inc., Framingham, MA); Escin; Digitonin; or *Gypsophila* or *Chenopodium quinoa* saponins. Other preferred formulations include more than one saponin in the adjuvant combinations of the present invention, for example combinations of at least two of the following group comprising QS21, QS7, Quil A, β -escin, or digitonin.

Alternatively the saponin formulations may be combined with vaccine vehicles composed of chitosan or other polycationic polymers, polylactide and polylactide-co-glycolide particles, poly-N-acetyl glucosamine-based polymer matrix, particles composed of polysaccharides or chemically modified polysaccharides, liposomes and lipid-based particles, particles composed of glycerol monoesters, etc. The saponins may also be formulated in the presence of cholesterol to form particulate structures such as liposomes or ISCOMs. Furthermore, the saponins may be formulated together with a polyoxyethylene ether or ester, in either a non-particulate solution or suspension, or in a particulate structure such as a paucilamellar liposome or ISCOM. The saponins may also be formulated with excipients such as Carbopol^R to increase viscosity, or may be formulated in a dry powder form with a powder excipient such as lactose.

In one preferred embodiment, the adjuvant system includes the combination of a monophosphoryl lipid A and a saponin derivative, such as the combination of QS21 and 3D-MPL[®] adjuvant, as described in WO 94/00153, or a less reactogenic composition where the QS21 is quenched with cholesterol, as described in WO 96/33739. Other preferred formulations comprise an oil-in-water emulsion and tocopherol. Another

particularly preferred adjuvant formulation employing QS21, 3D-MPL[®] adjuvant and tocopherol in an oil-in-water emulsion is described in WO 95/17210.

Another enhanced adjuvant system involves the combination of a CpG-containing
5 oligonucleotide and a saponin derivative particularly the combination of CpG and QS21 as disclosed in WO 00/09159. Preferably the formulation additionally comprises an oil in water emulsion and tocopherol.

Additional illustrative adjuvants for use in the pharmaceutical compositions of the
10 invention include Montanide ISA 720 (Seppic, France), SAF (Chiron, California, United States), ISCOMS (CSL), MF-59 (Chiron), the SBAS series of adjuvants (*e.g.*, SBAS-2 or SBAS-4, available from SmithKline Beecham, Rixensart, Belgium), **Detox (Enhanzyn[®])** (Corixa, Hamilton, MT), RC-529 (Corixa, Hamilton, MT) and other aminoalkyl glucosaminide 4-phosphates (AGPs), such as those described in pending U.S. Patent
15 Application Serial Nos. 08/853,826 and 09/074,720, the disclosures of which are incorporated herein by reference in their entireties, and polyoxyethylene ether adjuvants such as those described in WO 99/52549A1.

Other preferred adjuvants include adjuvant molecules of the general formula (I):



20 Wherein, *n* is 1-50, A is a bond or -C(O)-, R is C₁₋₅₀ alkyl or Phenyl C₁₋₅₀ alkyl.

One embodiment of the present invention consists of a vaccine formulation comprising a polyoxyethylene ether of general formula (I), wherein *n* is between 1 and 50, preferably 4-24, most preferably 9; the R component is C₁₋₅₀, preferably C₄-C₂₀ alkyl and most preferably C₁₂ alkyl, and A is a bond. The concentration of the polyoxyethylene ethers
25 should be in the range 0.1-20%, preferably from 0.1-10%, and most preferably in the range 0.1-1%. Preferred polyoxyethylene ethers are selected from the following group: polyoxyethylene-9-lauryl ether, polyoxyethylene-9-stearyl ether, polyoxyethylene-8-stearyl ether, polyoxyethylene-4-lauryl ether, polyoxyethylene-35-lauryl ether, and polyoxyethylene-23-lauryl ether. Polyoxyethylene ethers such as polyoxyethylene lauryl

ether are described in the Merck index (12th edition: entry 7717). These adjuvant molecules are described in WO 99/52549.

The polyoxyethylene ether according to the general formula (I) above may, if desired, be
5 combined with another adjuvant. For example, a preferred adjuvant combination is preferably with CpG as described in the pending UK patent application GB 9820956.2.

Preferably a carrier is also present in the vaccine composition according to the invention.
10 The carrier may be an oil in water emulsion, or an aluminium salt, such as aluminium phosphate or aluminium hydroxide.

A preferred oil-in-water emulsion comprises a metabolisable oil, such as squalene, alpha tocopherol and Tween 80. In a particularly preferred aspect the antigens in the vaccine
15 composition according to the invention are combined with QS21 and 3D-MPL in such an emulsion. Additionally the oil in water emulsion may contain span 85 and/or lecithin and/or tricaprylin.

Typically for human administration QS21 and 3D-MPL will be present in a vaccine in the
20 range of 1µg - 200µg, such as 10-100µg, preferably 10µg - 50µg per dose. Typically the oil in water will comprise from 2 to 10% squalene, from 2 to 10% alpha tocopherol and from 0.3 to 3% tween 80. Preferably the ratio of squalene: alpha tocopherol is equal to or less than 1 as this provides a more stable emulsion. Span 85 may also be present at a level of 1%. In some cases it may be advantageous that the vaccines of the present
25 invention will further contain a stabiliser.

Non-toxic oil in water emulsions preferably contain a non-toxic oil, e.g. squalane or squalene, an emulsifier, e.g. Tween 80, in an aqueous carrier. The aqueous carrier may be, for example, phosphate buffered saline.

30

A particularly potent adjuvant formulation involving QS21, 3D-MPL and tocopherol in an oil in water emulsion is described in WO 95/17210.

The present invention also provides a polyvalent vaccine composition comprising a vaccine
5 formulation of the invention in combination with other antigens, in particular antigens useful for treating cancers, autoimmune diseases and related conditions. Such a polyvalent vaccine composition may include a TH-1 inducing adjuvant as hereinbefore described.

According to another embodiment of this invention, an immunogenic composition
10 described herein is delivered to a host via antigen presenting cells (APCs), such as dendritic cells, macrophages, B cells, monocytes and other cells that may be engineered to be efficient APCs. Such cells may, but need not, be genetically modified to increase the capacity for presenting the antigen, to improve activation and/or maintenance of the T cell response, to have anti-tumor effects *per se* and/or to be immunologically compatible
15 with the receiver (*i.e.*, matched HLA haplotype). APCs may generally be isolated from any of a variety of biological fluids and organs, including tumor and peritumoral tissues, and may be autologous, allogeneic, syngeneic or xenogeneic cells.

Certain preferred embodiments of the present invention use dendritic cells or progenitors
20 thereof as antigen-presenting cells. Dendritic cells are highly potent APCs (Banchereau and Steinman, *Nature* 392:245-251, 1998) and have been shown to be effective as a physiological adjuvant for eliciting prophylactic or therapeutic antitumor immunity (*see* Timmerman and Levy, *Ann. Rev. Med.* 50:507-529, 1999). In general, dendritic cells may be identified based on their typical shape (stellate *in situ*, with marked cytoplasmic
25 processes (dendrites) visible *in vitro*), their ability to take up, process and present antigens with high efficiency and their ability to activate naïve T cell responses. Dendritic cells may, of course, be engineered to express specific cell-surface receptors or ligands that are not commonly found on dendritic cells *in vivo* or *ex vivo*, and such modified dendritic cells are contemplated by the present invention. As an alternative to
30 dendritic cells, secreted vesicles antigen-loaded dendritic cells (called exosomes) may be used within a vaccine (*see* Zitvogel et al., *Nature Med.* 4:594-600, 1998).

Dendritic cells and progenitors may be obtained from peripheral blood, bone marrow, tumor-infiltrating cells, peritumoral tissues-infiltrating cells, lymph nodes, spleen, skin, umbilical cord blood or any other suitable tissue or fluid. For example, dendritic cells may be differentiated *ex vivo* by adding a combination of cytokines such as GM-CSF, IL-4, IL-13 and/or TNF α to cultures of monocytes harvested from peripheral blood.

Alternatively, CD34 positive cells harvested from peripheral blood, umbilical cord blood or bone marrow may be differentiated into dendritic cells by adding to the culture medium combinations of GM-CSF, IL-3, TNF α , CD40 ligand, LPS, flt3 ligand and/or other compound(s) that induce differentiation, maturation and proliferation of dendritic cells.

Dendritic cells are conveniently categorized as "immature" and "mature" cells, which allows a simple way to discriminate between two well characterized phenotypes. However, this nomenclature should not be construed to exclude all possible intermediate stages of differentiation. Immature dendritic cells are characterized as APC with a high capacity for antigen uptake and processing, which correlates with the high expression of Fc γ receptor and mannose receptor. The mature phenotype is typically characterized by a lower expression of these markers, but a high expression of cell surface molecules responsible for T cell activation such as class I and class II MHC, adhesion molecules (*e.g.*, CD54 and CD11) and costimulatory molecules (*e.g.*, CD40, CD80, CD86 and 4-1BB).

APCs may generally be transfected with a polynucleotide of the invention (or portion or other variant thereof) such that the encoded polypeptide, or an immunogenic portion thereof, is expressed on the cell surface. Such transfection may take place *ex vivo*, and a pharmaceutical composition comprising such transfected cells may then be used for therapeutic purposes, as described herein. Alternatively, a gene delivery vehicle that targets a dendritic or other antigen presenting cell may be administered to a patient, resulting in transfection that occurs *in vivo*. *In vivo* and *ex vivo* transfection of dendritic cells, for example, may generally be performed using any methods known in the art, such as those described in WO 97/24447, or the gene gun approach described by Mahvi et al., *Immunology and cell Biology* 75:456-460, 1997. Antigen loading of dendritic cells may be achieved by incubating dendritic cells or progenitor cells with the tumor polypeptide,

DNA (naked or within a plasmid vector) or RNA; or with antigen-expressing recombinant bacterium or viruses (*e.g.*, vaccinia, fowlpox, adenovirus or lentivirus vectors). Prior to loading, the polypeptide may be covalently conjugated to an immunological partner that provides T cell help (*e.g.*, a carrier molecule). Alternatively,
5 a dendritic cell may be pulsed with a non-conjugated immunological partner, separately or in the presence of the polypeptide.

While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, the type of carrier will typically vary
10 depending on the mode of administration. Compositions of the present invention may be formulated for any appropriate manner of administration, including for example, topical, oral, nasal, mucosal, intravenous, intracranial, intraperitoneal, subcutaneous and intramuscular administration.

Carriers for use within such pharmaceutical compositions are biocompatible, and may
15 also be biodegradable. In certain embodiments, the formulation preferably provides a relatively constant level of active component release. In other embodiments, however, a more rapid rate of release immediately upon administration may be desired. The formulation of such compositions is well within the level of ordinary skill in the art using known techniques. Illustrative carriers useful in this regard include microparticles of
20 poly(lactide-co-glycolide), polyacrylate, latex, starch, cellulose, dextran and the like. Other illustrative delayed-release carriers include supramolecular biovectors, which comprise a non-liquid hydrophilic core (*e.g.*, a cross-linked polysaccharide or oligosaccharide) and, optionally, an external layer comprising an amphiphilic compound, such as a phospholipid (*see e.g.*, U.S. Patent No. 5,151,254 and PCT applications WO
25 94/20078, WO/94/23701 and WO 96/06638). The amount of active compound contained within a sustained release formulation depends upon the site of implantation, the rate and expected duration of release and the nature of the condition to be treated or prevented.

In another illustrative embodiment, biodegradable microspheres (*e.g.*, polylactate
30 polyglycolate) are employed as carriers for the compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4,897,268; 5,075,109; 5,928,647; 5,811,128; 5,820,883; 5,853,763; 5,814,344, 5,407,609 and

5,942,252. Modified hepatitis B core protein carrier systems. such as described in WO/99 40934, and references cited therein, will also be useful for many applications. Another illustrative carrier/delivery system employs a carrier comprising particulate-protein complexes, such as those described in U.S. Patent No. 5,928,647, which are
5 capable of inducing a class I-restricted cytotoxic T lymphocyte responses in a host.

The pharmaceutical compositions of the invention will often further comprise one or more buffers (*e.g.*, neutral buffered saline or phosphate buffered saline), carbohydrates (*e.g.*, glucose, mannose, sucrose or dextrans), mannitol, proteins, polypeptides or amino acids such as glycine, antioxidants, bacteriostats, chelating agents such as EDTA or
10 glutathione, adjuvants (*e.g.*, aluminum hydroxide), solutes that render the formulation isotonic, hypotonic or weakly hypertonic with the blood of a recipient, suspending agents, thickening agents and/or preservatives. Alternatively, compositions of the present invention may be formulated as a lyophilizate.

15 The pharmaceutical compositions described herein may be presented in unit-dose or multi-dose containers, such as sealed ampoules or vials. Such containers are typically sealed in such a way to preserve the sterility and stability of the formulation until use. In general, formulations may be stored as suspensions, solutions or emulsions in oily or aqueous vehicles. Alternatively, a pharmaceutical composition may be stored in a freeze-
20 dried condition requiring only the addition of a sterile liquid carrier immediately prior to use.

The development of suitable dosing and treatment regimens for using the particular compositions described herein in a variety of treatment regimens, including *e.g.*, oral,
25 parenteral, intravenous, intranasal, and intramuscular administration and formulation, is well known in the art, some of which are briefly discussed below for general purposes of illustration.

In certain applications, the pharmaceutical compositions disclosed herein may be
30 delivered *via* oral administration to an animal. As such, these compositions may be formulated with an inert diluent or with an assimilable edible carrier, or they may be

enclosed in hard- or soft-shell gelatin capsule, or they may be compressed into tablets, or they may be incorporated directly with the food of the diet.

The active compounds may even be incorporated with excipients and used in the form of ingestible tablets, buccal tables, troches, capsules, elixirs, suspensions, syrups, wafers, and the like (see, for example, Mathiowitz *et al.*, Nature 1997 Mar 27;386(6623):410-4; Hwang *et al.*, Crit Rev Ther Drug Carrier Syst 1998;15(3):243-84; U. S. Patent 5,641,515; U. S. Patent 5,580,579 and U. S. Patent 5,792,451). Tablets, troches, pills, capsules and the like may also contain any of a variety of additional components, for example, a binder, such as gum tragacanth, acacia, cornstarch, or gelatin; excipients, such as dicalcium phosphate; a disintegrating agent, such as corn starch, potato starch, alginic acid and the like; a lubricant, such as magnesium stearate; and a sweetening agent, such as sucrose, lactose or saccharin may be added or a flavoring agent, such as peppermint, oil of wintergreen, or cherry flavoring. When the dosage unit form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit. For instance, tablets, pills, or capsules may be coated with shellac, sugar, or both. Of course, any material used in preparing any dosage unit form should be pharmaceutically pure and substantially non-toxic in the amounts employed. In addition, the active compounds may be incorporated into sustained-release preparation and formulations.

Typically, these formulations will contain at least about 0.1% of the active compound or more, although the percentage of the active ingredient(s) may, of course, be varied and may conveniently be between about 1 or 2% and about 60% or 70% or more of the weight or volume of the total formulation. Naturally, the amount of active compound(s) in each therapeutically useful composition may be prepared in such a way that a suitable dosage will be obtained in any given unit dose of the compound. Factors such as solubility, bioavailability, biological half-life, route of administration, product shelf life, as well as other pharmacological considerations will be contemplated by one skilled in the art of preparing such pharmaceutical formulations, and as such, a variety of dosages and treatment regimens may be desirable.

For oral administration the compositions of the present invention may alternatively be incorporated with one or more excipients in the form of a mouthwash, dentifrice, buccal tablet, oral spray, or sublingual orally-administered formulation. Alternatively, the active
5 ingredient may be incorporated into an oral solution such as one containing sodium borate, glycerin and potassium bicarbonate, or dispersed in a dentifrice, or added in a therapeutically-effective amount to a composition that may include water, binders, abrasives, flavoring agents, foaming agents, and humectants. Alternatively the compositions may be fashioned into a tablet or solution form that may be placed under
10 the tongue or otherwise dissolved in the mouth.

In certain circumstances it will be desirable to deliver the pharmaceutical compositions disclosed herein parenterally, intravenously, intramuscularly, or even intraperitoneally. Such approaches are well known to the skilled artisan, some of which are further
15 described, for example, in U. S. Patent 5,543,158; U. S. Patent 5,641,515 and U. S. Patent 5,399,363. In certain embodiments, solutions of the active compounds as free base or pharmacologically acceptable salts may be prepared in water suitably mixed with a surfactant, such as hydroxypropylcellulose. Dispersions may also be prepared in glycerol, liquid polyethylene glycols, and mixtures thereof and in oils. Under ordinary
20 conditions of storage and use, these preparations generally will contain a preservative to prevent the growth of microorganisms.

Illustrative pharmaceutical forms suitable for injectable use include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation of sterile
25 injectable solutions or dispersions (for example, see U. S. Patent 5,466,468). In all cases the form must be sterile and must be fluid to the extent that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms, such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (*e.g.*, glycerol, propylene glycol, and liquid polyethylene glycol, and the like),
30 suitable mixtures thereof, and/or vegetable oils. Proper fluidity may be maintained, for

example, by the use of a coating, such as lecithin, by the maintenance of the required particle size in the case of dispersion and/or by the use of surfactants. The prevention of the action of microorganisms can be facilitated by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the
5 like. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride. Prolonged absorption of the injectable compositions can be brought about by the use in the compositions of agents delaying absorption, for example, aluminum monostearate and gelatin.

10 In one embodiment, for parenteral administration in an aqueous solution, the solution should be suitably buffered if necessary and the liquid diluent first rendered isotonic with sufficient saline or glucose. These particular aqueous solutions are especially suitable for intravenous, intramuscular, subcutaneous and intraperitoneal administration. In this connection, a sterile aqueous medium that can be employed will be known to those of
15 skill in the art in light of the present disclosure. For example, one dosage may be dissolved in 1 ml of isotonic NaCl solution and either added to 1000 ml of hypodermoclysis fluid or injected at the proposed site of infusion, (see for example, "Remington's Pharmaceutical Sciences" 15th Edition, pages 1035-1038 and 1570-1580). Some variation in dosage will necessarily occur depending on the condition of the subject
20 being treated. Moreover, for human administration, preparations will of course preferably meet sterility, pyrogenicity, and the general safety and purity standards as required by FDA Office of Biologics standards.

In another embodiment of the invention, the compositions disclosed herein may be
25 formulated in a neutral or salt form. Illustrative pharmaceutically-acceptable salts include the acid addition salts (formed with the free amino groups of the protein) and which are formed with inorganic acids such as, for example, hydrochloric or phosphoric acids, or such organic acids as acetic, oxalic, tartaric, mandelic, and the like. Salts formed with the free carboxyl groups can also be derived from inorganic bases such as, for example,
30 sodium, potassium, ammonium, calcium, or ferric hydroxides, and such organic bases as isopropylamine, trimethylamine, histidine, procaine and the like. Upon formulation,

solutions will be administered in a manner compatible with the dosage formulation and in such amount as is therapeutically effective.

The carriers can further comprise any and all solvents, dispersion media, vehicles,
5 coatings, diluents, antibacterial and antifungal agents, isotonic and absorption delaying agents, buffers, carrier solutions, suspensions, colloids, and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, its use in the therapeutic compositions is contemplated. Supplementary active ingredients
10 can also be incorporated into the compositions. The phrase "pharmaceutically-acceptable" refers to molecular entities and compositions that do not produce an allergic or similar untoward reaction when administered to a human.

In certain embodiments, the pharmaceutical compositions may be delivered by intranasal
15 sprays, inhalation, and/or other aerosol delivery vehicles. Methods for delivering genes, nucleic acids, and peptide compositions directly to the lungs *via* nasal aerosol sprays has been described, *e.g.*, in U. S. Patent 5,756,353 and U. S. Patent 5,804,212. Likewise, the delivery of drugs using intranasal microparticle resins (Takenaga *et al.*, J Controlled Release 1998 Mar 2;52(1-2):81-7) and lysophosphatidyl-glycerol compounds (U. S.
20 Patent 5,725,871) are also well-known in the pharmaceutical arts. Likewise, illustrative transmucosal drug delivery in the form of a polytetrafluoroethylene support matrix is described in U. S. Patent 5,780,045.

In certain embodiments, liposomes, nanocapsules, microparticles, lipid particles, vesicles,
25 and the like, are used for the introduction of the compositions of the present invention into suitable host cells/organisms. In particular, the compositions of the present invention may be formulated for delivery either encapsulated in a lipid particle, a liposome, a vesicle, a nanosphere, or a nanoparticle or the like. Alternatively, compositions of the present invention can be bound, either covalently or non-covalently, to the surface of
30 such carrier vehicles.

The formation and use of liposome and liposome-like preparations as potential drug carriers is generally known to those of skill in the art (see for example, Lasic, Trends Biotechnol 1998 Jul;16(7):307-21; Takakura, Nippon Rinsho 1998 Mar;56(3):691-5; Chandran *et al.*, Indian J Exp Biol. 1997 Aug;35(8):801-9; Margalit, Crit Rev Ther Drug
5 Carrier Syst. 1995;12(2-3):233-61; U.S. Patent 5,567,434; U.S. Patent 5,552,157; U.S. Patent 5,565,213; U.S. Patent 5,738,868 and U.S. Patent 5,795,587, each specifically incorporated herein by reference in its entirety).

Liposomes have been used successfully with a number of cell types that are normally
10 difficult to transfect by other procedures, including T cell suspensions, primary hepatocyte cultures and PC 12 cells (Renneisen *et al.*, J Biol Chem. 1990 Sep 25;265(27):16337-42; Muller *et al.*, DNA Cell Biol. 1990 Apr;9(3):221-9). In addition, liposomes are free of the DNA length constraints that are typical of viral-based delivery systems. Liposomes have been used effectively to introduce genes, various drugs,
15 radiotherapeutic agents, enzymes, viruses, transcription factors, allosteric effectors and the like, into a variety of cultured cell lines and animals. Furthermore, the use of liposomes does not appear to be associated with autoimmune responses or unacceptable toxicity after systemic delivery.

20 In certain embodiments, liposomes are formed from phospholipids that are dispersed in an aqueous medium and spontaneously form multilamellar concentric bilayer vesicles (also termed multilamellar vesicles (MLVs)).

Alternatively, in other embodiments, the invention provides for pharmaceutically-
25 acceptable nanocapsule formulations of the compositions of the present invention. Nanocapsules can generally entrap compounds in a stable and reproducible way (see, for example, Quintanar-Guerrero *et al.*, Drug Dev Ind Pharm. 1998 Dec;24(12):1113-28). To avoid side effects due to intracellular polymeric overloading, such ultrafine particles (sized around 0.1 μm) may be designed using polymers able to be degraded *in vivo*. Such
30 particles can be made as described, for example, by Couvreur *et al.*, Crit Rev Ther Drug Carrier Syst. 1988;5(1):1-20; zur Muhlen *et al.*, Eur J Pharm Biopharm. 1998

Mar;45(2):149-55; Zambaux *et al.* J Controlled Release. 1998 Jan 2;50(1-3):31-40; and U. S. Patent 5,145,684.

5 This invention also relates to the use of polynucleotides, in the form of primers derived from the polynucleotides of the present invention, and of polypeptides, in the form of antibodies or reagents specific for the polypeptide of the present invention, as diagnostic reagents.

10 The identification of genetic or biochemical markers in blood or tissues that will enable the detection of very early changes along the carcinogenesis pathway will help in determining the best treatment for the patient. Surrogate tumour markers, such as polynucleotide expression, can be used to diagnose different forms and states of cancer. The identification of expression levels of the polynucleotides of the invention will be useful in both the staging of the cancerous disorder and grading the nature of the cancerous tissue. The staging process monitors the advancement of the cancer and is determined on the presence or
15 absence of malignant tissue in the areas biopsied. The polynucleotides of the invention can help to perfect the staging process by identifying markers for the aggressivity of a cancer, for example the presence in different areas of the body. The grading of the cancer describes how closely a tumour resembles normal tissue of its same type and is assessed by its cell morphology and other markers of differentiation. The polynucleotides of the invention can
20 be useful in determining the tumour grade as they can help in the determination of the differentiation status of the cells of a tumour.

The diagnostic assays offer a process for diagnosing or determining a susceptibility to cancers, autoimmune disease and related conditions through diagnosis by methods
25 comprising determining from a sample derived from a subject an abnormally decreased or increased level of polypeptide or mRNA. This method of diagnosis is known as differential expression. The expression of a particular gene is compared between a diseased tissue and a normal tissue. A difference between the polynucleotide-related gene, mRNA, or protein in the two tissues is compared, for example in molecular weight,
30 amino acid or nucleotide sequence, or relative abundance, indicates a change in the gene, or a gene which regulates it, in the tissue of the human that was suspected of being diseased.

Decreased or increased expression can be measured at the RNA level. PolyA RNA is first isolated from the two tissues and the detection of mRNA encoded by a gene corresponding to a differentially expressed polynucleotide of the invention can be detected by, for example, in situ hybridization in tissue sections, reverse transcriptase-PCR, using Northern blots containing poly A+ mRNA, or any other direct or indirect RNA detection method. An increased or decreased expression of a given RNA in a diseased tissue compared to a normal tissue suggests that the transcript and/or the expressed protein has a role in the disease. Thus detection of a higher or lower level of mRNA corresponding to SEQ ID NO 1 or 3 relative to normal level is indicative of the presence of cancer in the patient.

mRNA expression levels in a sample can be determined by generation of a library of expressed sequence tags (ESTs) from the sample. The relative representation of ESTs in the library can be used to assess the relative representation of the gene transcript in the starting sample. The EST analysis of the test can then be compared to the EST analysis of a reference sample to determine the relative expression levels of the polynucleotide of interest.

Other mRNA analyses can be carried out using serial analysis of gene expression (SAGE) methodology (Velculescu et. Al. Science (1995) 270:484), differential display methodology (For example, US 5,776,683) or hybridization analysis which relies on the specificity of nucleotide interactions.

Alternatively, the comparison could be made at the protein level. The protein sizes in the two tissues may be compared using antibodies to detect polypeptides in Western blots of protein extracts from the two tissues. Expression levels and subcellular localization may also be detected immunologically using antibodies to the corresponding protein. Further assay techniques that can be used to determine levels of a protein, such as a polypeptide of the present invention, in a sample derived from a host are well-known to those of skill in the art. A raised or decreased level of polypeptide expression in the diseased tissue compared with the same protein expression level in the normal tissue indicates that the expressed protein may be involved in the disease.

In the assays of the present invention, the diagnosis can be determined by detection of gene product expression levels encoded by at least one sequence set forth in SEQ ID NOS: 1 or 3. A comparison of the mRNA or protein levels in a diseased versus normal tissue may also be
5 used to follow the progression or remission of a disease.

A large number of polynucleotide sequences in a sample can be assayed using polynucleotide arrays. These can be used to examine differential expression of genes and to determine gene function. For example, arrays of the polynucleotide sequences SEQ ID NO:
10 1 or 3 can be used to determine if any of the polynucleotides are differentially expressed between a normal and cancer cell. In one embodiment of the invention, an array of oligonucleotides probes comprising the SEQ ID NO: 1 or 3 nucleotide sequence or fragments thereof can be constructed to conduct efficient screening of e.g., genetic mutations. Array technology methods are well known and have general applicability and
15 can be used to address a variety of questions in molecular genetics including gene expression, genetic linkage, and genetic variability (see for example: M.Chee et al., Science, Vol 274, pp 610-613 (1996)).

“Diagnosis” as used herein includes determination of a subject’s susceptibility to a
20 disease, determination as to whether a subject presently has the disease, and also the prognosis of a subject affected by the disease.

The present invention, further relates to a diagnostic kit for performing a diagnostic assay which comprises:

- 25 (a) a polynucleotide of the present invention, preferably the nucleotide sequence of SEQ ID NO: 1 or 3, or a fragment thereof;
- (b) a nucleotide sequence complementary to that of (a);
- (c) a polypeptide of the present invention, preferably the polypeptide of SEQ ID NO: 2 or 4, or a fragment thereof; or
- 30 (d) an antibody to a polypeptide of the present invention, preferably to the polypeptide of SEQ ID NO:2 OR 4.

The nucleotide sequences of the present invention are also valuable for chromosomal localisation. The sequence is specifically targeted to, and can hybridize with, a particular location on an individual human chromosome. The mapping of relevant sequences to chromosomes according to the present invention is an important first step in correlating those sequences with gene associated disease. Once a sequence has been mapped to a precise chromosomal location, the physical position of the sequence on the chromosome can be correlated with genetic map data. Such data are found in, for example, V. McKusick, Mendelian Inheritance in Man (available on-line through Johns Hopkins University Welch Medical Library). The relationship between genes and diseases that have been mapped to the same chromosomal region are then identified through linkage analysis (coinheritance of physically adjacent genes). The differences in the cDNA or genomic sequence between affected and unaffected individuals can also be determined.

The polypeptides of the invention or their fragments or analogs thereof, or cells expressing them, can also be used as immunogens to produce antibodies immunospecific for polypeptides of the present invention. The term "immunospecific" means that the antibodies have substantially greater affinity for the polypeptides of the invention than their affinity for other related polypeptides in the prior art.

In a further aspect the invention provides an antibody immunospecific for a polypeptide according to the invention or an immunological fragment thereof as hereinbefore defined. Preferably the antibody is a monoclonal antibody

Antibodies generated against polypeptides of the present invention may be obtained by administering the polypeptides or epitope-bearing fragments, analogs or cells to an animal, preferably a non-human animal, using routine protocols. For preparation of monoclonal antibodies, any technique which provides antibodies produced by continuous cell line cultures can be used. Examples include the hybridoma technique (Kohler, G. and Milstein, C., Nature (1975) 256:495-497), the trioma technique, the human B-cell hybridoma technique (Kozbor *et al.*, Immunology Today (1983) 4:72) and the EBV-hybridoma technique (Cole *et al.*, Monoclonal Antibodies and Cancer Therapy, 77-96, Alan R. Liss, Inc., 1985).

Techniques for the production of single chain antibodies, such as those described in U.S. Patent No. 4,946,778, can also be adapted to produce single chain antibodies to polypeptides of this invention. Also, transgenic mice, or other organisms, including other mammals, may be used to express humanized antibodies.

5

The above-described antibodies may be employed to isolate or to identify clones expressing the polypeptide or to purify the polypeptides by affinity chromatography.

The antibody of the invention may also be employed to prevent or treat cancer, particularly ovarian and colon cancer, autoimmune disease and related conditions.

10

Another aspect of the invention relates to a method for inducing or modulating an immunological response in a mammal which comprises inoculating the mammal with a polypeptide of the present invention, adequate to produce antibody and/or T cell immune response to protect or ameliorate the symptoms or progression of the disease. Yet

15

another aspect of the invention relates to a method of inducing or modulating immunological response in a mammal which comprises, delivering a polypeptide of the present invention *via* a vector directing expression of the polynucleotide and coding for the polypeptide *in vivo* in order to induce such an immunological response to produce antibody to protect said animal from diseases.

20

It will be appreciated that the present invention therefore provides a method of treating abnormal conditions such as, for instance, cancer and autoimmune diseases, in particular, ovarian and colon cancer, related to either a presence of, an excess of, or an under-expression of, CASB6411 polypeptide activity.

25

The present invention further provides for a method of screening compounds to identify those which stimulate or which inhibit the function of the CASB6411 polypeptide. In general, agonists or antagonists may be employed for therapeutic and prophylactic purposes for such diseases as hereinbefore mentioned. Compounds may be identified from a variety of sources, for example, cells, cell-free preparations, chemical libraries, and natural product mixtures. Such agonists, antagonists or inhibitors so-identified may be natural or modified substrates, ligands, receptors, enzymes, etc., as the case may be, of the polypeptide; or may be structural or functional mimetics thereof (see Coligan *et al.*, Current Protocols in

30

Immunology 1(2):Chapter 5 (1991)). Screening methods will be known to those skilled in the art. Further screening methods may be found in for example D. Bennett *et al.*, J Mol Recognition, 8:52-58 (1995); and K. Johanson *et al.*, J Biol Chem, 270(16):9459-9471 (1995) and references therein.

5

Thus the invention provides a method for screening to identify compounds which stimulate or which inhibit the function of the polypeptide of the invention which comprises a method selected from the group consisting of:

- 10 (a) measuring the binding of a candidate compound to the polypeptide (or to the cells or membranes bearing the polypeptide) or a fusion protein thereof by means of a label directly or indirectly associated with the candidate compound;
- (b) measuring the binding of a candidate compound to the polypeptide (or to the cells or membranes bearing the polypeptide) or a fusion protein thereof in the presense of a labeled competitor;
- 15 (c) testing whether the candidate compound results in a signal generated by activation or inhibition of the polypeptide, using detection systems appropriate to the cells or cell membranes bearing the polypeptide;
- (d) mixing a candidate compound with a solution containing a polypeptide of claim 1, to form a mixture, measuring activity of the polypeptide in the mixture, and comparing the
- 20 activity of the mixture to a standard; or
- (e) detecting the effect of a candidate compound on the production of mRNA encoding said polypeptide and said polypeptide in cells, using for instance, an ELISA assay.

The polypeptide of the invention may be used to identify membrane bound or soluble
25 receptors, if any, through standard receptor binding techniques known in the art. Well known screening methods may also be used to identify agonists and antagonists of the polypeptide of the invention which compete with the binding of the polypeptide of the invention to its receptors, if any.

30 Thus, in another aspect, the present invention relates to a screening kit for identifying agonists, antagonists, ligands, receptors, substrates, enzymes, etc. for polypeptides of the present invention; or compounds which decrease or enhance the production of such polypeptides, which comprises:

- (a) a polypeptide of the present invention;
 - (b) a recombinant cell expressing a polypeptide of the present invention;
 - (c) a cell membrane expressing a polypeptide of the present invention; or
 - (d) antibody to a polypeptide of the present invention;
- 5 which polypeptide is preferably that of SEQ ID NO:2 OR 4.

It will be readily appreciated by the skilled artisan that a polypeptide of the present invention may also be used in a method for the structure-based design of an agonist, antagonist or inhibitor of the polypeptide, by:

- 10 (a) determining in the first instance the three-dimensional structure of the polypeptide;
- (b) deducing the three-dimensional structure for the likely reactive or binding site(s) of an agonist, antagonist or inhibitor;
- (c) synthesizing candidate compounds that are predicted to bind to or react with the deduced binding or reactive site; and
- 15 (d) testing whether the candidate compounds are indeed agonists, antagonists or inhibitors.

Gene therapy may also be employed to effect the endogenous production of CASB6411 polypeptide by the relevant cells in the subject. For an overview of gene therapy, see

20 Chapter 20, Gene Therapy and other Molecular Genetic-based Therapeutic Approaches, (and references cited therein) in Human Molecular Genetics, T Strachan and A P Read, BIOS Scientific Publishers Ltd (1996).

Vaccine preparation is generally described in Pharmaceutical Biotechnology, Vol.61

25 Vaccine Design - the subunit and adjuvant approach, edited by Powell and Newman, Plenum Press, 1995. New Trends and Developments in Vaccines, edited by Voller et al., University Park Press, Baltimore, Maryland, U.S.A. 1978. Encapsulation within liposomes is described, for example, by Fullerton, U.S. Patent 4,235,877. Conjugation of proteins to macromolecules is disclosed, for example, by Likhite, U.S. Patent 4,372,945

30 and by Armor et al., U.S. Patent 4,474,757.

The amount of protein in each vaccine dose is selected as an amount which induces an immunoprotective response without significant, adverse side effects in typical vaccinees.

Such amount will vary depending upon which specific immunogen is employed. Generally, it is expected that each dose will comprise 1-1000 μ g of protein, preferably 2-100 μ g, most preferably 4-40 μ g. An optimal amount for a particular vaccine can be ascertained by standard studies involving observation of antibody titres and other
5 responses in subjects. Following an initial vaccination, subjects may receive a boost in about 4 weeks.

"Isolated" means altered "by the hand of man" from the natural state. If an "isolated" composition or substance occurs in nature, it has been changed or removed from its
10 original environment, or both. For example, a polynucleotide or a polypeptide naturally present in a living animal is not "isolated," but the same polynucleotide or polypeptide separated from the coexisting materials of its natural state is "isolated", as the term is employed herein.

15 "Polynucleotide" generally refers to any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA including single and double stranded regions.

"Variant" refers to a polynucleotide or polypeptide that differs from a reference
20 polynucleotide or polypeptide, but retains essential properties. A typical variant of a polynucleotide differs in nucleotide sequence from another, reference polynucleotide. Changes in the nucleotide sequence of the variant may or may not alter the amino acid sequence of a polypeptide encoded by the reference polynucleotide. Nucleotide changes may result in amino acid substitutions, additions, deletions, fusions and truncations in the
25 polypeptide encoded by the reference sequence, as discussed below. A typical variant of a polypeptide differs in amino acid sequence from another, reference polypeptide. Generally, differences are limited so that the sequences of the reference polypeptide and the variant are closely similar overall and, in many regions, identical. A variant and reference polypeptide may differ in amino acid sequence by one or more substitutions,
30 additions, deletions in any combination. A substituted or inserted amino acid residue may or may not be one encoded by the genetic code. A variant of a polynucleotide or polypeptide may be a naturally occurring such as an allelic variant, or it may be a variant

that is not known to occur naturally. Non-naturally occurring variants of polynucleotides and polypeptides may be made by mutagenesis techniques or by direct synthesis.

"Identity," as known in the art, is a relationship between two or more polypeptide sequences or two or more polynucleotide sequences, as determined by comparing the sequences. In the art, "identity" also means the degree of sequence relatedness between polypeptide or polynucleotide sequences, as the case may be, as determined by the match between strings of such sequences. "Identity" and "similarity" can be readily calculated by known methods, including but not limited to those described in (Computational Molecular Biology, Lesk, A.M., ed., Oxford University Press, New York, 1988; Biocomputing: Informatics and Genome Projects, Smith, D.W., ed., Academic Press, New York, 1993; Computer Analysis of Sequence Data, Part I, Griffin, A.M., and Griffin, H.G., eds., Humana Press, New Jersey, 1994; Sequence Analysis in Molecular Biology, von Heinje, G., Academic Press, 1987; and Sequence Analysis Primer, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991; and Carillo, H., and Lipman, D., SIAM *J. Applied Math.*, 48: 1073 (1988). Preferred methods to determine identity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in publicly available computer programs. Preferred computer program methods to determine identity and similarity between two sequences include, but are not limited to, the GCG program package (Devereux, J., et al., *Nucleic Acids Research* 12(1): 387 (1984)), BLASTP, BLASTN, and FASTA (Atschul, S.F. et al., *J. Molec. Biol.* 215: 403-410 (1990). The BLAST X program is publicly available from NCBI and other sources (BLAST Manual, Altschul, S., et al., NCBI NLM NIH Bethesda, MD 20894; Altschul, S., et al., *J. Mol. Biol.* 215: 403-410 (1990). The well known Smith Waterman algorithm may also be used to determine identity.

The preferred algorithm used is FASTA. The preferred parameters for polypeptide or polynucleotide sequence comparison using this algorithm include the following:

Gap Penalty:12

Gap extension penalty: 4

Word size: 2, max 6

Preferred parameters for polypeptide sequence comparison with other methods include the following:

1) Algorithm: Needleman and Wunsch, J. Mol Biol. 48: 443-453 (1970)

Comparison matrix: BLOSSUM62 from Hentikoff and Hentikoff, Proc. Natl. Acad. Sci.

5 USA. 89:10915-10919 (1992)

Gap Penalty: 12

Gap Length Penalty: 4

A program useful with these parameters is publicly available as the "gap" program from
10 Genetics Computer Group, Madison WI. The aforementioned parameters are the default parameters for polypeptide comparisons (along with no penalty for end gaps).

Preferred parameters for polynucleotide comparison include the following:

1) Algorithm: Needleman and Wunsch, J. Mol Biol. 48: 443-453 (1970)

15 Comparison matrix: matches = +10, mismatch = 0

Gap Penalty: 50

Gap Length Penalty: 3

A program useful with these parameters is publicly available as the "gap" program from
20 Genetics Computer Group, Madison WI. The aforementioned parameters are the default parameters for polynucleotide comparisons.

By way of example, a polynucleotide sequence of the present invention may be identical to the reference sequence of SEQ ID NO:1, that is be 100% identical, or it may include
25 up to a certain integer number of nucleotide alterations as compared to the reference sequence. Such alterations are selected from the group consisting of at least one nucleotide deletion, substitution, including transition and transversion, or insertion, and wherein said alterations may occur at the 5' or 3' terminal positions of the reference nucleotide sequence or anywhere between those terminal positions, interspersed either
30 individually among the nucleotides in the reference sequence or in one or more contiguous groups within the reference sequence. The number of nucleotide alterations is determined by multiplying the total number of nucleotides in SEQ ID NO:1 by the

numerical percent of the respective percent identity (divided by 100) and subtracting that product from said total number of nucleotides in SEQ ID NO:1, or:

$$n_n \leq x_n - (x_n \cdot y),$$

wherein n_n is the number of nucleotide alterations, x_n is the total number of nucleotides in SEQ ID NO:1, and y is, for instance, 0.70 for 70%, 0.80 for 80%, 0.85 for 85%, 0.90 for 90%, 0.95 for 95%, etc., and wherein any non-integer product of x_n and y is rounded down to the nearest integer prior to subtracting it from x_n . Alterations of a polynucleotide sequence encoding the polypeptide of SEQ ID NO:2 OR 4 may create nonsense, missense or frameshift mutations in this coding sequence and thereby alter the polypeptide encoded by the polynucleotide following such alterations.

Similarly, a polypeptide sequence of the present invention may be identical to the reference sequence of SEQ ID NO:2 OR 4, that is be 100% identical, or it may include up to a certain integer number of amino acid alterations as compared to the reference sequence such that the % identity is less than 100%. Such alterations are selected from the group consisting of at least one amino acid deletion, substitution, including conservative and non-conservative substitution, or insertion, and wherein said alterations may occur at the amino- or carboxy-terminal positions of the reference polypeptide sequence or anywhere between those terminal positions, interspersed either individually among the amino acids in the reference sequence or in one or more contiguous groups within the reference sequence. The number of amino acid alterations for a given % identity is determined by multiplying the total number of amino acids in SEQ ID NO:2 OR 4 by the numerical percent of the respective percent identity (divided by 100) and then subtracting that product from said total number of amino acids in SEQ ID NO:2 OR 4, or:

$$n_a \leq x_a - (x_a \cdot y),$$

wherein n_a is the number of amino acid alterations, x_a is the total number of amino acids in SEQ ID NO:2 OR 4, and y is, for instance 0.70 for 70%, 0.80 for 80%, 0.85 for 85% etc., and wherein any non-integer product of x_a and y is rounded down to the nearest integer prior to subtracting it from x_a .

“Homolog” is a generic term used in the art to indicate a polynucleotide or polypeptide sequence possessing a high degree of sequence relatedness to a subject sequence. Such relatedness may be quantified by determining the degree of identity and/or similarity between the sequences being compared as hereinbefore described. Falling within this generic term are the terms “ortholog”, meaning a polynucleotide or polypeptide that is the functional equivalent of a polynucleotide or polypeptide in another species and “paralog” meaning a functionally similar sequence when considered within the same species.

10 Examples

Example 1

Subtractive cDNA cloning of colon tumour-associated antigen (TAA) candidates.

Subtractive cDNA libraries are produced using standard technologies. Briefly, total RNA is extracted from frozen (-70°C) tumour and matched normal colon samples using the TriPure reagent and protocol (Boehringer). Target RNA is prepared by pooling total RNA from three tumour samples (30 µg each). Driver RNA is prepared by pooling total RNA from three matched normal colon samples (10 µg each) and total RNA from seven normal tissues other than colon (brain, heart, kidney, liver, bladder, skin, spleen; 10 µg each). Total RNA from non-colon normal tissues is purchased from InVitrogen.

Messenger RNA is purified from total RNA using oligo-dT magnetic bead technology (Dynal) and quantified by spectrofluorimetry (BioRad).

Target and driver mRNA are reverse transcribed into cDNA using one of two strategies: 1) Target sequences for PCR oligonucleotides are introduced onto the ends of the newly synthesised cDNA during reverse transcription using the template switching capability of reverse transcriptase (ClonTech SMART PCR cDNA synthesis kit). 2) Alternatively, the target and driver mRNA are reverse transcribed into cDNA using an oligo-dT primer and converted to double-strand cDNA; the cDNA is cleaved with RsaI and linkers for PCR amplification are ligated onto the extremities of the cDNA fragments.

In both cases, target and driver cDNA are amplified by long range PCR (ClonTech SMART PCR Synthesis Kit and Advantage PCR Polymerase Mix) and used as starting material for subtractive cloning. For amplification, cycling conditions and optimisation of the number of PCR cycles are as described in the Advantage PCR protocol.

Two subtractive cloning strategies are used: ClonTech PCR SELECT (see ClonTech kit protocol and N. Gurskaya *et al.* 1996. Analytical Biochemistry: 240, 90) and cRDA (M. Hubank and D. Schatz. 1994. Nucleic Acids Research: 22, 5640). When the PCR SELECT protocol is used, the primary PCR SELECT subtraction products are submitted to a supplementary round of cRDA subtraction. When the cRDA protocol is used, two consecutive cycles of cRDA subtraction are performed. In each case the products of both cycles of subtraction are cloned into pCR-TOPO (Invitrogen) and transformed into *E. coli* to produce a subtracted cDNA plasmid library.

An alternative strategy is also followed: subtraction of normal colon sequences and sequences from non-colon normal tissues are subtracted in separate hybridizations. In this case, target and driver RNA are assembled for the first subtraction as above with the exception that non-colon RNA is left out of the driver pool and amounts of normal colon are increased to 10 µg. Preparation of target and driver cDNA and subtractive hybridization are performed as described above. A second subtraction is then performed on the products of the first subtraction, but the driver is now composed of a pool of normal colon and normal non-colon mRNA from the seven normal tissues.

Example 2

Differential Screening of cDNA arrays.

Identification of tumour-associated genes in the subtracted cDNA library is accomplished by differential screening.

Total bacterial DNA is extracted from 100 µl over-night cultures. Bacteria are lysed with guanidium isothiocyanate and the bacterial DNA is affinity purified using magnetic glass (Boehringer). Plasmid inserts are recovered from the bacterial DNA by Advantage PCR amplification (Clontech). The PCR products are dotted onto two nylon membranes to produce high density cDNA arrays using the Biomek 96 HDRT tool (Beckman). The spotted cDNA is covalently linked to the membrane by UV irradiation. The first membrane is hybridised with a mixed cDNA probe prepared from the tumour of a single patient. The second membrane is hybridised with an equivalent amount of mixed cDNA probe prepared from normal colon of the same patient. The probe cDNA is prepared by PCR amplification as described above and is labelled using the AlkPhos Direct System (Amersham).

Hybridisation conditions and stringency washes are as described in the AlkPhos Direct kit. Hybridized probe is detected by chemiluminescence. Hybridisation intensities for each cDNA fragment on both blots are measured by film densitometry or direct measurement (BioRad Fluor-S Max). The ratio of the tumour to normal hybridisation intensities (T/N) is calculated for each gene to evaluate the degree of over-expression in the tumour. Genes which are significantly over-expressed in colon tumours are followed-up. Significance is arbitrarily defined as one standard deviation of the T/N frequency distribution. Differential screening experiments are repeated using RNA from multiple patient donors (>18) to estimate the frequency of over-expressing tumours in the patient population.

In addition, the DNA arrays are hybridised with mixed cDNA probes from normal tissues other than colon (see list above) to determine the level of expression of the candidate gene in these tissues.

Example 3

Real-time RT-PCR analysis

Real-time RT-PCR (U. Gibson. 1996. Genome Research: 6,996) is used to compare mRNA transcript abundance of the candidate antigen in matched tumour and normal colon tissues from multiple patients. In addition, mRNA levels of the candidate gene in a panel of normal tissues are also evaluated by this approach.

Total RNA from normal and tumour colon is extracted from snap frozen biopsies using TriPure reagent (Boehringer). Total RNA from normal tissues is purchased from InVitrogen or is extracted from snap frozen biopsies using TriPure reagent. Poly-A+ mRNA is purified from total RNA after DNAase treatment using oligo-dT magnetic beads (Dyna). Quantification of the mRNA is performed by spectrofluorimetry (VersaFluor, BioRad) using SybrII dye (Molecular Probes). Primers for real-time PCR amplification are designed with the Perkin-Elmer Primer Express software using default options for TaqMan amplification conditions.

Real-time reactions are assembled according to standard PCR protocols using 2 ng of purified mRNA for each reaction. SybrI dye (Molecular Probes) is added at a final dilution of 1/75000 for real-time detection. Amplification (40 cycles) and real-time detection is performed in a Perkin-Elmer Biosystems PE7700 system using conventional

instrument settings. Ct values are calculated using the PE7700 Sequence Detector software. Several Ct values are obtained for each samples : for the patient samples, the tumour Ct (CtT) and the matched normal colon Ct (CtN) values on the candidate TAA, and for the panel of normal tissue samples, a CtXY for each normal tissue XY. An
 5 another Ct (CtA) is also calculated on Actin gene, as an internal reference, for all of the samples.

As the efficiency of PCR amplification under the prevailing experimental conditions is close to the theoretical amplification efficiency, $2^{(CtN/T/XY-CtA)}$ value is an estimate of the
 10 relative TAA transcript level of the sample, standardised with respect to Actin transcript level. A value of 1 thus suggests the candidate antigen and Actin have the same expression level.

Real-time PCR reactions were performed on tumour colon and adjacent normal colon
 15 from biopsies of 18 patients. Results are shown in figure 1. 36 normal tissue samples, representing 28 different tissues (see table 2), were also tested by the same procedure. Results are shown in figure 2.

TAA transcript levels are calculated as described above. The proportion of patients over-
 20 expressing the candidate antigen, as well as the average transcript over-expression versus normal tissues is also calculated from this data set.

Overall results are shown in Table 1 :

Table 1 : CASB6411 Real-time PCR expression results

25

% of patients with a CASB6411 transcript level higher in tumour colon than adjacent normal colon (positive patients)	83%
% of positive patients with a CASB6411 transcript level at least 3 fold higher CASB6411 transcript than adjacent normal colon	60%
% of positive patients with a CASB6411 transcript level at least 10 fold higher in tumour colon than adjacent normal colon	40%
Average transcript over-expression fold in positive patients	40
% of patients with a CASB6411 transcript level higher in tumour colon than normal tissue median	100%
% of patients with a mRNA level at least 3 fold higher in tumour colon than normal tissue median	100%

% of patients with a mRNA level at least 10 fold higher in tumour colon than normal tissue median	40%
Normal tissues where CASB6411 transcript expression is equivalent than tumour transcript level in tumours	Lung, prostate

Table 1 clearly suggest CASB6411 transcript is over-expressed in colorectal tumours compared to adjacent normal colon and to most of the above mentioned normal tissues.

- 5 More than 80% of the patients strongly over-express CASB6411 transcript in tumour, as compared to adjacent normal colon. Average over-expression fold is at least of 40, with 40 % of patients having overexpressing at least 10 fold. More over, all of the patients over-express CASB6411 transcript in colorectal tumors, as compared to other normal tissues, 40 % of them overexpressing it at least 10 fold.

10

Table 2 : listing of normal tissues used for CASB7439 transcript expression analysis.

Tissue	Abbreviation	Categorie
Adrenal gland	Ad_Gl	non dispensable
Aorta	Ao	non dispensable
Bladder	Bl	non dispensable
Bone marrow	Bo_Ma	non dispensable
Brain	Bra	non dispensable
Cervix	Ce	non dispensable
Colon	Co	non dispensable
Fallopian tube	Fa_Tu	non dispensable
Heart	He	non dispensable
Ileum	Il	non dispensable
Kidney	Ki	non dispensable
Liver	Li	non dispensable
Lung	Lu	non dispensable
Lymph node	Ly_No	non dispensable
Oesophagus	Oe	non dispensable
Parathyroid gland	Pa_Thy	non dispensable
Rectum	Re	non dispensable
Skin	Sk	non dispensable
Skeletal muscle	Sk_Mu	non dispensable
Small intestine	Sm_In	non dispensable
Spleen	Sp	non dispensable
Stomach	St	non dispensable
Thyroid gland	Thy	non dispensable
Trachea	Tra	non dispensable
Ovary	Ov	dispensable
Placenta	Pl	dispensable
Prostate	Pr	dispensable
Testis	Te	dispensable

Example 4

DNA microarrays

DNA micro-arrays are used to examine mRNA expression profiles of large collections of genes in multiple samples. This information is used to complement the data obtained by
5 real-time PCR and provides an independent measure of gene expression levels in tumors and normal tissues.

Examples of current technologies for production of DNA micro-arrays include 1) The Affymetrix "GeneChip" arrays in which oligonucleotides are synthesized on the surface
10 of the chip by solid phase chemical synthesis using a photolithographic process 2) DNA spotting technology in which small volumes of a DNA solution are robotically deposited and then immobilized onto the surface of a solid phase (e.g. glass). In both instances, the chips are hybridized with cDNA or cRNA which has been extracted from the tissue of interest (e.g. normal tissue, tumour etc...) and labeled with radioactivity or with a
15 fluorescent reporter molecule. The labeled material is hybridized to the chip and the amount of probe bound to each sequence on the chip is determined using a specialized scanner. The experiment can be set-up with a single fluorescent reporter (or radioactivity) or, alternatively, can be performed using two fluorescent reporters. In this latter case, each of the two samples is labeled with one of the reporter molecules. The two labeled
20 samples are then hybridized competitively to the sequences on the DNA chip. The ratio of the two fluorescent signals is determined for each sequence on the chip. This ratio is used to calculate the relative abundance of the transcript in the two samples. Detailed protocols are available from a number of sources including "DNA Microarrays: A practical approach. Schena M. Oxford University Press 1999" and the World Wide Web
25 (<http://cmgm.stanford.edu/pbrown/protocols/index.html>), <http://arrayit.com/DNA-Microarray-Protocols/>) and specialized distributors (e.g. Affymetrix).

Example 5

Northern-Southern blot analysis

Limited amounts of mixed tumour and matched normal colon cDNA are amplified by Advantage PCR (see above). Messenger RNA from multiple normal tissues is also amplified using the same procedure. The amplified cDNA (1 µg) is electrophoresed on a

1.2% agarose gel and transferred onto a nylon membrane. The membrane is hybridised (AlkPhos Direct System) with a probe prepared using a fragment of the candidate TAA cDNA. Northern-Southern analysis provides information on transcript size, presence of splice variants and transcript abundance in tumour and normal tissues.

5

Example 6

Northern Blot Analysis

Northern blots are produced according to standard protocols using 1 µg of poly A+ mRNA. Radioactive probes are prepared using the Ready-to-Go system (Pharmacia).

10

Example 7

In silico detection of the full length cDNA sequence

EST sequence databases are screened with experimentally obtained cDNA sequence fragments, using the Blast algorithm (Altschul, S.F., Gish, W., Miller, W., Myers, E.W. & Lipman, D.J. (1990) "Basic local alignment search tool." J. Mol. Biol. 215:403-41).

15

The aim is to search for overlapping or longer identical EST sequences. Matched EST sequences are then assembled together, using the SeqMan software of the Lasergene package (DNASTAR). The consensus sequence of the resulting assembly is an EST-derived longer cDNA. This EST-derived cDNA is further analysed using the GeneMark software to find a potential open reading frame (ORF). The translated sequence of the ORF is compared with protein databases, using the Blast algorithm, to find homologues. If any, the homologous protein sequences are further used to complete the cDNA prediction by searching for genomic contig homologies using the Wise2 algorithm, leading to a genome-derived, virtual cDNA sequence. This virtual cDNA is finally assembled with EST-derived cDNA, and the new consensus cDNA undergoes a final check against ESTs to confirm the Wise2 prediction, and correct potential sequencing errors and frameshifts. The virtual cDNA is considered as a virtual full length cDNA once a full ORF (from start to stop codons), with clear protein homologies and coding potential.

20

25

30

The SEQ ID NO:1 has been obtained using this *in silico* cloning method of the full length cDNA sequence of CASB6411, which has a putative open reading frame of 460 amino acids (SEQ ID NO:2), and a potential isoform generated by alternative splicing (SEQ ID NO:3), encoding a truncated protein (SEQ ID NO:4).

Virtual full-length cDNA sequence is experimentally checked as described below.

Example 8

Experimental Identification of the full length cDNA sequence

- 5 Colon tumour cDNA libraries are constructed using the Lambda Zap II system (Stratagene) from 5 µg of polyA+ mRNA. The supplied protocol is followed except that SuperscriptII (Life Technologies) is used for the reverse transcription step. Oligo dT-primed and random-primed libraries are constructed. About 1.5×10^6 independent phages are plated for each screening of the library. Phage plaques are transferred onto nylon
- 10 filters and hybridised using a cDNA probe labelled with AlkPhos Direct. Positive phages are detected by chemiluminescence. Positive phage are excised from the agar plat, eluted in 500µl SM buffer and confirmed by gene-specific PCR. Eluted phages are converted to single strand M13 bacteriophage by in vivo excision. The bacteriophage is then converted to double strand plasmid DNA by infection of E. coli. Infected bacteria are plated and
- 15 submitted to a second round of screening with the cDNA probe. Plasmid DNA is purified from positive bacterial clones and sequenced on both strands.

- When the full length gene cannot be obtained directly from the cDNA library, missing sequence is isolated using RACE technology (Marathon Kit, ClonTech.). This approach
- 20 relies on reverse transcribing mRNA into double strand cDNA, ligating linkers onto the ends of the cDNA and amplifying the desired extremity of the cDNA using a gene-specific primer and one of the linker oligonucleotides. Marathon PCR products are cloned into a plasmid (pCRII-TOPO, InVitrogen) and sequenced.

25 Example 9.

EST profiles

- A complementary approach to experimental antigen tissue expression characterization is to explore the human EST database. ESTs ('Expressed Sequence Tags') are small fragments of cDNA made from a collection of mRNA extracted from a particular tissue
- 30 or cell line. Such database currently provides a massive amount of human ESTs (10^6) from several hundreds of cDNA tissue libraries, including tumoral tissues from various types and states of disease. By means of informatics tools (Blast), a comparison search of

the CASB6411 sequence is performed in order to have further insight into tissue expression.

EST distribution of CASB6411 :

EST GenBank Accession number	EST cDNA tissue library
C00562	Human adult (K.Okubo)
C05837	Human pancreatic islet
AA172076	Stratagene ovarian cancer (#937219)
AA172244	Stratagene ovarian cancer (#937219)
AA612697	NCI_CGAP_Co10
AA371314	Prostate gland I
AI264367	NCI_CGAP_Co8
AI270207	NCI_CGAP_Co14
AI278830	NCI_CGAP_Co8
AI281230	NCI_CGAP_Co8
AI283827	NCI_CGAP_Co8
AI285194	NCI_CGAP_Co8
AI285227	NCI_CGAP_Co8
AI346622	NCI_CGAP_Co8
AI362363	NCI_CGAP_Gas4
AI473464	NCI_CGAP_Gas4
AI697014	NCI_CGAP_Pan1
AI799626	NCI_CGAP_Gas4
AI830044	NCI_CGAP_Lu19
AI921465	NCI_CGAP_Gas4
AI925050	NCI_CGAP_Gas4
AW029127	NCI_CGAP_Gas4
AW365013	DT0057
AW452356	NCI_CGAP_Sub5
AW469177	NCI_CGAP_Gas4
AW469181	NCI_CGAP_Gas4
AW582253	ST0212
AW810203	ST0125
AW810268	ST0125
AW810418	ST0125
AW814058	ST0198
AW869793	SN0075

5

SEQ ID NO:1 perfectly aligns with 32 ESTs : 13 are from 2 stomach cDNA libraries, 9 are from 3 tumor colon libraries, 2 are from one tumor ovary library, one is from one tumor pancreas library, one is from one lung tumor library, one is from one mixed tumor library, one is from one normal prostate library, one is from one normal stomach library, one is from normal pancreas library, 2 are from libraries of unknown type.

10

This clearly suggests, as expected, that CASB6411 is over-expressed in tumor tissues, with an emphasis in colorectal and stomach tumor tissues, as compared to normal tissues.

5 Example 10 :

10.1 Expression and purification of tumour-specific antigens

Expression in microbial hosts, or alternatively in vitro transcription/translation, is used to produce the antigen of the invention for vaccine purposes and to produce protein
10 fragments or whole protein for rapid purification and generation of antibodies needed for characterization of the naturally expressed protein by immunohistochemistry or for follow-up of purification.

Recombinant proteins may be expressed in two microbial hosts, *E. coli* and in yeast (such as *Saccharomyces cerevisiae* or *Pichia pastoris*). This allows the selection of the
15 expression system with the best features for this particular antigen production. In general, the recombinant antigen will be expressed in *E. coli* and the reagent protein expressed in yeast.

The expression strategy first involves the design of the primary structure of the recombinant antigen. In general an expression fusion partner (EFP) is placed at the N
20 terminal extremity to improve levels of expression that could also include a region useful for modulating the immunogenic properties of the antigen, an immune fusion partner (IFP). In addition, an affinity fusion partner (AFP) useful for facilitating further purification is included at the C-terminal end.

When the recombinant strains are available, the recombinant product is characterized by
25 the evaluation of the level of expression and the prediction of further solubility of the protein by analysis of the behavior in the crude extract.

After growth on appropriate culture medium and induction of the recombinant protein expression, total extracts are analyzed by SDS-PAGE. The recombinant proteins are visualized in stained gels and identified by Western blot analysis using specific
30 antibodies.

A comparative evaluation of the different versions of the expressed antigen will allow the selection of the most promising candidate that is to be used for further purification and immunological evaluation.

The purification scheme follows a classical approach based on the presence of an His affinity tail in the recombinant protein. In a typical experiment the disrupted cells are filtered and the acellular extracts loaded onto an Ion Metal Affinity Chromatography (IMAC; Ni⁺⁺NTA from Qiagen) that will specifically retain the recombinant protein.
5 The retained proteins are eluted by 0-500 mM Imidazole gradient (possibly in presence of a detergent) in a phosphate buffer.

10.2 Antibody production and immunohistochemistry

- 10 Small amounts of relatively purified protein can be used to generate immunological tools in order to
- a) detect the expression by immunohistochemistry in normal or cancer tissue sections;
 - b) detect the expression, and to follow the protein during the purification process (ELISA/ Western Blot); or
 - 15 c) characterise/ quantify the purified protein (ELISA).

10.2.1 Polyclonal antibodies:

Immunization

- Rabbits are immunised , intramuscularly (I.M.) , 3 times at 3 weeks intervals with
20 100µg of protein, formulated in the adjuvant 3D-MPL/QS21. Three weeks after each immunisation a blood sample is taken and the antibody titer estimated in the serum by ELISA using the protein as coating antigen following a standard protocol.

ELISA

- 25 96 well microplates (maxisorb Nunc) are coated with 5µg of protein overnight at 4°C. After 1 hour saturation at 37°C with PBS NCS 1%, serial dilution of the rabbit sera is added for 1H 30 at 37°C (starting at 1/10). After 3 washings in PBS Tween, anti rabbit biotinylated anti serum (Amersham) is added (1/5000). Plates are washed and peroxidase coupled streptavidin (1/5000) is added for 30 min at 37°C. After washing,
30 50µl TMB (BioRad) is added for 7 min and the reaction then stopped with H2SO4 0.2M. The OD can be measured at 450 nm and midpoint dilutions calculated by SoftmaxPro.

10.2.2 Monoclonal antibodies:

Immunization

5 BALB/c mice are immunized 3 times at 3 week intervals with 5 µg of purified protein. Bleedings are performed 14 days post II and 1 week post 3. The sera are tested by Elisa on purified protein used as coated antigen. Based on these results (midpoint dilution > 10000) one mouse is selected for fusion

Fusion/ HATselection

Spleen cells are fused with the SP2/0 myeloma according to a standard protocol using PEG 40% and DMSO 5%. Cells are then seeded in 96 well plates $2.5 \times 10^4 - 10^5$ cells/well and resistant clones will be selected in HAT medium. The supernatant of these hybridomas will be tested for their content in specific antibodies and when positive, will be submitted to 2 cycles of limited dilution . After 2 rounds of screening, 3 hybridomas will be chosen for ascitis production.

15 10.2.3 Immunohistochemistry

When antibodies are available, immuno staining is performed on normal or cancer tissue sections, in order to determine :

- ◇ the level of expression of the antigen of the invention in cancer relative to normal tissue or
- 20 ◇ the proportion of cancer of a certain type expressing the antigen
- ◇ if other cancer types also express the antigen
- ◇ the proportion of cells expressing the antigen in a cancer tissue

25 Tissue sample preparation

After dissection, the tissue sample is mounted on a cork disk in OCT compound and rapidly frozen in isopentane previously super cooled in liquid nitrogen (-160°C). The block will then be conserved at -70°C until use. 7-10µm sections will be realised in a cryostat chamber (-20, -30°C).

30

Staining

Tissue sections are dried for 5 min at room Temperature (RT), fixed in acetone for 10min at RT, dried again, and saturated with PBS 0.5% BSA 5% serum. After 30 min at

RT either a direct or indirect staining is performed using antigen specific antibodies. A direct staining leads to a better specificity but a less intense staining whilst an indirect staining leads to a more intense but less specific staining.

5 10.3 Analysis of human cellular immune responses to the antigen of the invention

The immunological relevance of the antigen of the invention can be assessed by in vitro priming of human T cells. All T cell lymphocyte lines and dendritic cells are derived from PBMCs (peripheral blood mononuclear cells) of healthy donors (preferred HLA-A2 subtype). An HLA-A2.1/Kb transgenic mouse model is also used for screening of HLA-A2.1 peptides.

Newly discovered antigen-specific CD8+ T cell lines are raised and maintained by weekly in vitro stimulation. The lytic activity and the γ -IFN production of the CD8 lines in response to the antigen or antigen derived-peptides is tested using standard procedures.

Two strategies to raise the CD8+ T cell lines are used: a peptide-based approach and a whole gene-based approach. Both approaches require the full-length cDNA of the newly discovered antigen in the correct reading frame to be either cloned in an appropriate delivery system or to be used to predict the sequence of HLA binding peptides.

Peptide-based approach

Briefly, transgenic mice are immunized with adjuvanted HLA-A2 peptides, those unable to induce a CD8 response (as defined by an efficient lysis of peptide-pulsed autologous spleen cells) will be further analyzed in the human system.

Human dendritic cells (cultured according to Romani et al.) will be pulsed with peptides and used to stimulate CD8-sorted T cells (by FACS). After several weekly stimulations, the CD8 lines will be first tested on peptide-pulsed autologous BLCL (EBV-B transformed cell lines). To verify the proper in vivo processing of the peptide, the CD8 lines will be tested on cDNA-transfected tumour cells (HLA-A2 transfected LnCaP, Skov3 or CAMA tumour cells).

Whole gene-based approach

CD8+ T cell lines will be primed and stimulated with either gene-gun transfected dendritic cells, retrovirally transduced B7.1-transfected fibroblasts, recombinant pox virus (Kim et al.) or adenovirus (Butterfield et al.) infected dendritic cells. Virus infected cells are very efficient to present antigenic peptides since the antigen is expressed at high level but can only be used once to avoid the over-growth of viral T cells lines.

After alternated stimulations, the CD8+ lines are tested on cDNA-transfected tumour cells as indicated above. Peptide specificity and identity is determined to confirm the immunological validation.

CD4+ T-cell response

Similarly, the CD4+ T-cell immune response can also be assessed. Generation of specific CD4+ T-cells is made using dendritic cells loaded with recombinant purified protein or peptides to stimulate the T-cells.

Predicted epitopes (nonamers and decamers) binding HLA alleles :

The HLA Class I binding peptide sequences are predicted either by the Parker's algorithm (Parker, K. C., M. A. Bednarek, and J. E. Coligan. 1994. Scheme for ranking potential HLA-A2 binding peptides based on independent binding of individual peptide side-chains. J. Immunol. 152:163 and http://bimas.dcrf.nih.gov/molbio/hla_bind/) or the Rammensee method (Rammensee, Friede, Stevanovic, MHC ligands and peptide motifs: 1st listing, Immunogenetics 41, 178-228, 1995 ; Rammensee, Bachmann, Stevanovic: MHC ligands and peptide motifs. Landes Bioscience 1997, and <http://134.2.96.221/scripts/hlaserver.dll/home.htm>). Peptides are then screened in the HLA-A2.1/Kb transgenic mice model (Vitiello et al.).

The HLA Class II binding peptide sequences are predicted using the Tepitope algorithm, with a score cut-off set to 6 (Sturniolo, Hammer et al., Nature Biotechnology. 1999. 17:555-561).

The following tables gather the Class I and II predicted epitope sequences :

HLA-A1 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	379	ITEGRKIMIR	112.500	SEQ ID NO:9

HLA-A0201 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	191	LLMDFVFSL	27926.980	SEQ ID NO:10
2	292	FLLFFPSFT	2650.811	SEQ ID NO:11
3	285	QMMTFFIFL	2266.849	SEQ ID NO:12
4	286	MMTFFIFLL	1329.564	SEQ ID NO:13
5	403	FLIEKLIKL	926.658	SEQ ID NO:14
6	147	VLLIRNIFL	739.032	SEQ ID NO:15
7	240	LVWIGIFFC	450.023	SEQ ID NO:16
8	305	TLAITIWRRL	368.501	SEQ ID NO:17
9	22	LIFCWDFTV	348.892	SEQ ID NO:18
10	203	FLGEFLRRI	343.941	SEQ ID NO:19
11	112	LLLPFVVSC	273.114	SEQ ID NO:20
12	66	LLTRFSAYM	176.513	SEQ ID NO:21
13	21	KLIFCWDFT	119.495	SEQ ID NO:22

5

HLA-A0201 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	190	LLMDFVFSL	6811.458	SEQ ID NO:23
2	191	LLMDFVFSLV	3407.985	SEQ ID NO:24
3	285	QMMTFFIFLL	2893.757	SEQ ID NO:25
4	93	YLAEYNLEFL	2497.344	SEQ ID NO:26
5	21	KLIFCWDFTV	1411.906	SEQ ID NO:27
6	293	LLFFPSFTGV	831.216	SEQ ID NO:28
7	65	QLTRFSAYM	384.175	SEQ ID NO:29
8	239	TLVWIGIFFC	364.502	SEQ ID NO:30
9	284	SQMMTFFIFL	318.231	SEQ ID NO:31
10	111	VLLLPFVVSC	273.114	SEQ ID NO:32
11	162	ILCYWLNNTV	271.948	SEQ ID NO:33
12	366	TLIVLIITYL	270.234	SEQ ID NO:34
13	146	YVLLIRNIFL	174.977	SEQ ID NO:35
14	195	FVFSLVNSFL	174.977	SEQ ID NO:36
15	112	LLLPFVVSCI	150.931	SEQ ID NO:37
16	355	LIGSVHFFFI	147.808	SEQ ID NO:38
17	138	YEMPRHEVYV	126.598	SEQ ID NO:39
18	66	LLTRFSAYMV	118.238	SEQ ID NO:40

HLA A205 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	191	LLMDFVFSL	514.080	SEQ ID NO:10
2	403	FLIEKLIKL	126.000	SEQ ID NO:14

10

HLA A205 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID
1	190	LLLMDEVFSL	171.360	SEQ ID NO:23
2	146	YVLLIRNIFL	126.000	SEQ ID NO:35
3	195	FVFSLVNSFL	126.000	SEQ ID NO:36
4	285	QMMTFFIFLL	100.800	SEQ ID NO:25

HLA A24 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	14	IYSGGITKL	220.000	SEQ ID NO:41
2	91	VYYLAEYNL	200.000	SEQ ID NO:42
3	165	YYWLNTVAL	200.000	SEQ ID NO:43
4	187	IYRLLLMDF	120.000	SEQ ID NO:44

HLA A24 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score	SEQ ID :
1	145	VYVLLIRNIF	252.000	SEQ ID NO:45
2	164	CYYWLNTVAL	200.000	SEQ ID NO:46
3	92	YYLAEYNLEF	165.000	SEQ ID NO:47

HLA A3 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Parler's Score *	SEQ ID :
1	258	MLFIMFYSK	900.000	SEQ ID NO:48
2	148	LLIRNIFLK	135.000	SEQ ID NO:49
3	221	GLQEFDIAR	108.000	SEQ ID NO:50

- : Estimate of Half Time of Disassociation of a Molecule Containing This Subsequence.

HLA A3 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	401	KMFLIEKLIK	600.000	SEQ ID NO:51
2	257	IMLFIMFYSK	270.000	SEQ ID NO:52
3	407	KLIKLDMEK	180.000	SEQ ID NO:53
4	147	VLLIRNIFLK	135.000	SEQ ID NO:54

HLA B7 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	140	MPRHEVYVL	800.000	SEQ ID NO:55

HLA B7 : decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :

1	140	MPRHEVYVLL	800.000	SEQ ID NO:56
---	-----	------------	---------	--------------

HLA B4403 nonamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	138	YEMPRHEVY	480.000	SEQ ID NO:57

5

HLA B4403 decamers				
Rank	Start Position	Subsequence Residue Listing	Parker's Score *	SEQ ID :
1	179	WETLIGQDIY	120.000	SEQ ID NO:58

*: Estimate of half time of disassociation of a molecule containing this subsequence.

HLA-DRB1*1501 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Tepitope Score	SEQ ID :
1	363	VHFFILTLL	6.5	SEQ ID NO:59
2	210	LRRIIGMQL	6.2	SEQ ID NO:60

10

HLA-DRB1*0301 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Tepitope Score	SEQ ID :
1	193	LLMDFVFSLL	6.4	SEQ ID NO:61

HLA-DRB1*0703 : nonamers				
Rank	Start Position	Subsequence Residue Listing	Tepitope Score	SEQ ID :
1	216	MQLITSLGL	8.5	SEQ ID NO:62
2	197	FVFSLVNSF	8	SEQ ID NO:63
3	367	FILTLIVLI	8	SEQ ID NO:64
4	210	LRRIIGMQL	7.6	SEQ ID NO:60
5	381	WQITEGRKI	7.6	SEQ ID NO:65
6	148	VLLIRNIFL	7.3	SEQ ID NO:49
7	265	MFYSKNISL	7.3	SEQ ID NO:66
8	263	FIMFYSKNI	7.2	SEQ ID NO:67
9	305	VLCTLAITI	7.2	SEQ ID NO:17
10	284	WRASQMMTF	6.8	SEQ ID NO:68
11	365	FFFILTLIV	6.8	SEQ ID NO:69
12	133	FRLVERYEM	6.7	SEQ ID NO:70
13	366	FFILTLIVL	6.4	SEQ ID NO:71
14	238	YAQTLVWI	6.4	SEQ ID NO:72

SEQUENCE INFORMATION

SEQ ID NO:1

TGGGGAGGCAGAAGGCAGACTGATCACTTGAGGCCAGGAGTTTGAGACCTCAT
5 GTCTAAAAAAAAAAATTCTGTGAGGTGAGTTTTATTGTTATTCCCTCTCTACAG
ATATGGAACTGAGGCTGAGAATCAGAACCATTACAAAGACAAAAATCCCCAG
TTGGCAGATCCAGGGTTGCAAGCCAGGCCTGTGCAGCCCCAAAACAGTGCTTG
TTTAACCACTGTGTGGTGACCACACCGCTCCAGGCCAACAGCTTGGGGCTAAGT
CTTCACGTTGCCTTTCACCATTAAATAATAGGGCTGCCCTTTGTTGAAGCCCTGC
10 ACTCCCAGTGACGGCCATAATAACCTTCAGGTGTTCTGCTTTCTGCCTTCTCTAG
Catggccaagtatttcggaacaacttcattaatccccacatttactcggaggatcaccaagctgatctttgtgggacttactg
tcactcatgaaaagctgtgaagctaaaacagaagaatcttagcactgagataagggagaacctgtcagagctccgtcaggagaa
ttccaagttgacgttcaatcagctgctgacccgcttctgcctacatggttagcctgggtgtctctacaggagtggccatagcctgctg
tgcagccgtttattacctggctgagtacaacttagagttcctgaagacacacagtaacctggggcggtgctgttactgcttctgtgt
15 gtctgcattaatctggccgtgcatgcatctactccatgttcaggcttgaggaggtacgagatgccacggcacgaagtctacgttc
tctgatccgaaacatcttttgaaaatatcaatcattggcattcttgttactattggctcaacacggtggccctgtctggggaagagt
ttgggaaacctcattggccaggacatctaccgctccttctgattttgtgtctcttagtcaattccttctggggagtttctga
ggagaatcattgggatgcaactgatcacaagcttggccttcaggagtttgacattgccaggaaccttctagaactgatctatgcaca
aactctggtgtggattggcatcttcttctgccccctgctgccctttatccaatgattatgctttcatcatgttctactccaaaatatca
20 gcctgatgatgaatttcagcctccgagcaaaagctggcgccctcacagatgatgactttctcatcttctgtcttttcccatcctt
caccgggtcttgtgcacctggccatcaccatctggagattgaagccttcagctgactgtggcccttttcgaggtctgctctctcat
tactccatctacagctggatcgacacctaaagtacacggcctggtacctgtgggtgtttggatctatcggaacctcattggaagt
tgcactctttttcatctcaccctcattgtgctaatacatcattcttactggcagatcacagagggaaggaagattatgataaggc
tgctccatgagcagatcattaatgagggaagataaaatgttctgatagaaaaattgatcaagctgcaggatattggagaagaaa
25 gcaaacccagctcacttgttctggaaaggagaggtggagcaacaaggcttttgcatttgggggaacatgatggcagcttggac
ttgcgatctagaagatcagttcaagaagtaatccaaggcctgaTGACTCTTTTGGTAACCAGACACCAA
TCAAATAAGGGGAGGAGACGAAAATGGAATGATTTCTTCCATGCCACCTGTGCC
TTTAGGAACTGCCCAGAAGAAAATCCAAGGCTTTAGCCAGGAGCGGAAAACCTGAC
TACCATGTAATTATCAAAGTAAAATTGGGCATTCCATGCTATTTTTAATACCTGG
30 ATTGCTGATTTTTCAAGACAAAATACCTGGGGTTTTCCAATAAAGATTGTTGTAA
TATTGAAATGAGCCTACAAAAACCTAGGAAGAGATAACTAGGGAATAATGTATA
TTATCTTCAAGAAATGTGTGCAGGAATGATTGGTTCTTAGAAATCTCTCCTGCCA
GACTTCCCAGACCTGGCAAAGGTTTAGAACTGTTGCTAAGAAAAGTGGTCCAT
CCTGAATAAACATGTAATACTCCAGCAGGGATATGAAGCCTCTGAATTGTAGAA
35 CCTGCATTTATTTGTGACTTTGAACTAAAGACATCCCCCATGTCCCAAAGGTGG
AATAACAACCAGAGGTCTCATCTCTGAACCTTTCTTGCGTACTGATTACATGAGTCT

TTGGAGTCGGGGATGGAGGAGGTTCTGCCCCCTGTGAGGTGTTATACATGACCAT
CAAAGTCCTACGTCAAGCT

5 SEQ ID NO:2

MAKYFRNNFINPHIYSGGITKLIFCWDFTVTHEKAVKLKQKNLSTEIRENLSELRQENSKLTFNQLLTRFSA
YMWAVVSTGVAIACCAVYYLAEYNLEFLKTHSNPGAVLLLPFVVSCLAVPCIYSMFRLVERYEMPRHE
VYVLLIRNIFLKISIIIGILCYWLNTVALSGEECWETLIGQDIYRLLMDFVFSLVNSFLGEFLRRIIGMQL
10 ITSLGLQEFDIARNVLELIYAQTLVWIGIFFCPLLPFIQMIMLFIMFYSKNISLMMNFQPPSKAWRASQMMT
FFIFLLFFPSFTGVLCTLAITWRLKPSADCGPFRGLPLFIHSIYSWIDTLSTRPGYLWVWVIYRNIGSVH
FFFILTLIVLIITYLYWQITEGRKIMIRLLHEQIINEGKDKMFLIEKLIKLDMEKKANPSSVLRRREVEQ
QGFLHLGEHDGSLDLRSRRSVQEGNPRA

15 SEQ ID NO:3

TGGGGAGGCAGAAGGCAGACTGATCACTTGAGGCCAGGAGTTTGAGACCTCAT
GTCTAAAAAAAAAAATTCTGTGAGGTGAGTTTTATTGTIATTCCTCTCTACAG
ATATGGAAACTGAGGCTGAGAATCAGAACCATTACAAGACAAAAATCCCCCAG
20 TTGGCAGATCCAGGGTTGCAAGCCAGGCCTGTGCAGCCCCAAAACCAGTGCTTG
TTTAACCACTGTGTGGTGACCACACCGCTCCAGGCCAACAGCTTGGGGCTAAGT
CTTCACGTTGCCTTTCACCATTAATAATAGGGCTGCCCTTTGTTGAAGCCCTGC
ACTCCCAGTGACGGCCATAATAACCTTCAGGTGTTCTGCTTTCTGCCTTCTCTAG
Catggccaagtatttcggaacaacttcattaatccccacattactcggaggatcaccaagctgatctttgtctgggacttcactg
25 tcactcatgaaaaagctgtaagctaaaacagaagaatcttagcactgagataagggagaacctgtcagagctccgtcaggagaa
ttccaagttgacgttcaatcagctgctgacccgcttctctgctacatgtagcctgggtgtctctacaggagtggccatgacctgctg
tgcagccgtttattacctggctgagtacaacttaggttctgaagacacacagtaacctggggcggtgctgttactgctttctgtgt
gtctgcattctggccgtgcatctactccatgttcaggctgtggagaggtacgagatgccacggcacgaagtctacgttctcc
tgatccgcaggggattgatgtagTTCTCAAGTATGGGATGTACAGATGGGCAGGCAGTGAC
30 GCACAAAGGCTCCTGGGCTGAGGACGGGACTGAAATCATCCAGCGTTCCCCTTA
GTCAAGCTAAACATCTTTTTGAAAATATCAATCATTGGCATTCTTTGTTACTATT
GGCTCAACACCGTGGCCCTGTCTGGTGAAGAGTGTTGGGAAACCCCTCATTGGCC
AGGACATCTACCGGCTCCTTCTGATGGATTTTGTGTTCTCTTTAGTCAATTCCTT
CCTGGGGGAGTTTCTGAGGAGAATCATTGGGATGCAACTGATCACAAGTCTTGG
35 CCTTCAGGAGTTTGACATTGCCAGGAACGTTCTAGAACTGATCTATGCACAAAC
TCTGGTGTGGATTGGCATCTTCTTCTGCCCCCTGCTGCCCTTTATCCAAATGATT
ATGCTTTTCATCATGTTCTACTCCAAAAATATCAGCCTGATGATGAATTTCCAGC

CTCCGAGCAAAGCCTGGCGGGCCTCACAGATGATGACTTTCTTCATCTTCTTGC
TCTTTTCCCATCCTTCACCGGGGTCTTGTGCACCCTGGCCATCACCATCTGGA
GATTGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCCTCTCTTCATTC
ACTCCATCTACAGCTGGATCGACACCCTAAGTACACGGCCTGGCTACCTGTGGG
5 TTGTTTGGATCTATCGGAACCTCATTGGAAGTGTGCACTTCTTTTTTCATCCTCAC
CCTCATTGTGCTAATCATCACCTATCTTTACTGGCAGATCACAGAGGGAAGGAA
GATTATGATAAGGCTGCTCCATGAGCAGATCATTAAATGAGGGCAAAGATAAAAT
GTTCTGATAGAAAAATTGATCAAGCTGCAGGATATGGAGAAGAAAGCAAACCC
CAGCTCACTTGTTCTGGAAAGGAGAGAGGTGGAGCAACAAGGCTTTTTGCATTT
10 GGGGGAACATGATGGCAGTCTTGACTTGCGATCTAGAAGATCAGTTCAAGAAG
GTAATCCAAGGGCCTGATGACTCTTTTGGTAACCAGACACCAATCAAATAAGGG
GAGGAGACGAAAAATGGAATGATTTCTTCCATGCCACCTGTGCCTTTAGGAACTG
CCCAGAAGAAAATCCAAGGCTTTAGCCAGGAGCGGAAACTGACTACCATGTAAT
TATCAAAGTAAAATTGGGCATTCCATGCTATTTTAAATACCTGGATTGCTGATTT
15 TTCAAGACAAAATACTTGGGGTTTTCCAATAAAGATTGTTGTAATATTGAAATGA
GCCTACAAAACCTAGGAAGAGATAACTAGGGAATAATGTATATTATCTTCAAG
AAATGTGTGCAGGAATGATTGGTTCTTAGAAATCTCTCCTGCCAGACTTCCAG
ACCTGGCAAAGGTTTAGAACTGTTGCTAAGAAAAGTGGTCCATCCTGAATAAA
CATGTAATACTCCAGCAGGGATATGAAGCCTCTGAATTGTAGAACCTGCATTTA
20 TTTGTGACTTTGAACTAAAGACATCCCCCATGTCCCAAAGGTGGAATACAACCA
GAGGTCTCATCTCTGAACTTTCTTGCCTACTGATTACATGAGTCTTTGGAGTCG
GGGATGGAGGAGGTTCTGCCCCTGTGAGGTGTTATACATGACCATCAAAGTCCT
ACGTCAAGCT

SEQ ID NO:4

25

MAKYFRNNFINPHIYSGGITKLIFCWDFTVTHEKAVKLKQKNLSTEIRENLSELRQE
NSKLTFNQLLTRFSAYMVAWVSTGVAIACCAAVYYLAEYNLEFLKTHSNPGAVLL
LPFVVSCILAVPCIYSMFRLVERYEMPRHEVYVLLIRRGLM

30 SEQ ID NO:5

atcttttgctgggacttcactgtcactcatgaaaaagctgtgaagctaaaacagaagaatcttagcactgag
ataagggagaacctgtcagagctccgtcaggagaattccaagttgacgttcaatcagctgctgacccgcttc
tctgcctacatggttagcctgggttgctctacaggagtggccatagcctgctgtgcagccgtttattacctg
35 gctgagtacaacttagagttcctgaagacacagtaacctggggcggtgctgttactgcctttcggttg
tcctgcattaatctggcgtgccatgcatctactccatgttcaggcttggtggagaggtacgagatgccacgg

cacgaagtctacgttctcctgatccgaaacatctttttgaaaatatcaatcattggcattctttgttactat
 tggctcaacaccgtggccctgtctggtgaagagtgttgggaaaccctcattggccaggacatctaccggctc
 cttctgatggattttgtgttctctttagtcaattccttctctggggagtttctgaggagaatcattgggatg
 caactgatcacaagtcttggccttcaggagtttgacattgccaggaaacgttctagaactgatctatgcacaa
 5 actctggtgtggattggcatcttcttctgccccctgctgccctttatccaaatgattatgcttttcatcatg
 ttctactccaaaaatatcagcctgatgatgaatttccagcctccgagcaaagcctggcgggcctcacagatg
 atgactttcttcatcttcttcttcttccatctttcaccggggtcttgtgcaccctggccatcaccatc
 tggagattgaagccttcagctgactgtggcccttttccaggtctgcctctcttcattcactccatctacagc
 tggatcgacaccctaagtacacggcctggctacctgtgggttgttggatctatcggaacctcattggaagt
 10 gtgcacttcttttcatcctcaccctcattgtgctgatcatcacctatctttactggcagatcacagagggga
 aggaagattatgataaggctgctccatgagcagatcattaatgagggcaaagataaaatgttctctgatagaa
 aaattgatcaagctgcaggatattggagaagaaagcaaaccctcagctcacttgttctggaaggagagaggtg
 gagcaacaaggctttttgcatttgggggaacatgatggcagctcttgacttgcgatctagaagatcagttcaa
 gaaggtaatccaaggcctgaTGACTCTTTTGGTAACCAGACACCAATCAAATAAGGGGAGGAGATGAAAAT
 15 GGAATGATTTCTTCCATGCCACCTGTGCCTTTAGGAAGTCCCAGAAGAAAATCCAAGGCTTTAGCCAGGAG
 CGGAAACTGACTACCATGTAATTATCAAAGTAAAATTGGGCATTCCATGCTATTTTAAATACCTGGATTGCT
 GATTTTCAAGACAAAATACTTGGGGTTTTCCAATAAAGATTGTTGTAATATTGAAATGAGCCTACAAAAAC
 CTAGGAAGAGATAACTAGGAATAATGTATATTATCTTCAAGAAATGTGTGCAGGAATGATTGGTTCTTAGA
 AATCTCTCTGCCAGACTTCCCAGACCTGGCAAAGGTTTAGAAACTGTTGCTAAGAAAAGTGGTCCATCCTG
 20 AATAACATGTAATACTCCAGCAGGGATATGAAGCCTCTGAATTGTAGAACCTGCATTTATTTGTGACTTTG
 AACTAAGACATCCCCCATGTCCCAAAGGTGGAATACAACCAGAGGTCTCATCTCTGAACCTTCTTGCCTAC
 TGATTACATGAGTCTTTGGAGTCGGGGATGGAGGAGGTTCTGCCCTGTGAGGTGTTATACATGACCATCAA
 AGTCCTACGTCAAGCT

25 SEQ ID NO:6

IFCWDFVTTHEKAVKLKQKNLSTEIRENLSELRQENSKLTFNQLLTRFSAYMVAVVSTGVAIACCAAVYYL
 AEYNLEFLKTHSNPGAVLLLPFVVSINLAVPCISYMFRLVERYEMPRHEVYVLLIRNIFLKISIIIGILCY
 WLNTVALSGEECWETLIGQDIYRLLLMDFVFLVNSFLGEFLRRIIGMQLITSLGLQEFDIARNVLELIYAQ
 30 TLVWIGIFFCPLLPFIQMIMLFIMFYSKNISLMMNFQPPSKAWRASQMMTFFIFLLFFPSFTGVLCTLAITI
 WRLKPSADCGPFRGLPLFIHSIYSWIDTLSTRPGYLWVWVIYRNLIQSVHFFIFILTIVLIITYLYWQITEG
 RKIMIRLLHEQIINEGDKMFLIEKLIKLDMEKKANPSSVLRLERREVEQQGFLHLGEHDSLDLRSRRSVQ
 EGNPRA

35 SEQ ID NO:7

ctgatgatgaatttccagcctccgagcaaagcctggcgggcctcacagatgatgactttcttcatcttcttcttcttccatcttccac
 cggggcttctgtgacccctggccatcaccatctggagattgaagccttcagctgactgtggcccttttccaggtctgcctcttcttcatca
 ctccatctacagctggatcgacaccctaagtacacggcctggctacctgtgggttgttggatctatcggaacctcattggaagtgtgc

acttcttttctcctcaccctcattgtgtgatcatcacctatctttactggcagatcacagaggggaaggaagattatgataaggctgc
 tccatgagcagatcattaatgagggcaagataaaatgttctctgatagaaaattgatcaagctgcaggatatggagaagaaagc
 aaaccccagctcacttgttctgaaaggagagaggtggagcaacaaggcttttgcatttgggggaacatgatggcagcttctgactt
 gcatctagaagatcagttcaagaaggtaatccaaggcctgaTGA CTCTTTTGGTAACCAGACACCAAT
 5 CAAATAAGGGGAGGAGATGAAAATGGAATGATTTCTTCCATGCCACCTGTGCCT
 TTAGGAAGTGGCAGAAAGAAAATCCAAGGCTTTAGCCAGGAGCGGAAACTGACT
 ACCATGTAATTATCAAAGTAAAATTGGGCATTCCATGCTATTTTTTAATACCTGGA
 TTGCTGATTTTTCAAGACAAAATACTTGGGGTTTTCCAATAAAGATTGTTGTAAT
 ATTGAAATGAGCCTACAAAAACCTAGGAAGAGATAACTAGGGAATAATGTATAT
 10 TATCTTCAAGAAATGTGTGCAGGAATGATTGGTTCTTAGAAATCTCTCCTGCCA
 GACTTCCCAGACCTGGCAAAGGTTTAGAAACTGTTGCTAAGAAAAGTGGTCCAT
 CCTGAATAAACATGTAATACTCCAGCAGGGATATGAAGCCTCTGAATTGTAGAA
 CCTGCATTTATTTGTGACTTTGAACTAAAGACATCCCCCATGTCCCAAAGGTGG
 AATACAACCAGAGGTCTCATCTCTGAACTTTCTTGCGTACTGATTACATGAGTCT
 15 TTGGAGTCGGGGATGGAGGAGGTTCTGCCCCTGTGAGGTGTTATACATGACCAT
 CAAAGTCCTACGTCAAGCT

SEQ ID NO:8

20 LMMNFQPPSKAWRASQMMTFFIFLLFFPSFTGVLCTLAITIWRLKPSADCGPFRGLPLFIHSIYSWIDTLST
 RPGYLWVWVIYRNIGSVHFFILTLIVLIITYLYWQITEGRKIMIRLLHEQIINEGKDKMFLIEKLIKLOD
 MEKKANPSSLVLERREVEQQGFLHLGEHDGSLDLRSRRSVQEGNPRA

SEQ ID NO:9

25 ITEGRKIMIR

SEQ ID NO:10

30 LLMDFVFSL

SEQ ID NO:11

35 FLLFFPSFT

SEQ ID NO:12

40 QMMTFFIFL

SEQ ID NO:13

45 MMTFFIFLL

SEQ ID NO:14

FLIEKLIK
SEQ ID NO:15
5 VLLIRNIFL
SEQ ID NO:16
LVWIGIFFC
10 SEQ ID NO:17
TLAITIWRL
15 SEQ ID NO:18
LIFCWDFTV
SEQ ID NO:19
20 FLGEFLRRI
SEQ ID NO:20
25 LLLPFVWSC
SEQ ID NO:21
LLTRFSAYM
30 SEQ ID NO:22
KLIFCWDF
35 SEQ ID NO:23
LLMDFVFS
SEQ ID NO:24
40 LLMDFVFSV
SEQ ID NO:25
45 QMTFFIFLL
SEQ ID NO:26
YLAEYNLEFL
50 SEQ ID NO:27
KLIFCWDFTV
55 SEQ ID NO:28
LLFFPSFTGV

SEQ ID NO:29

QLLTRFSAYM

5 **SEQ ID NO:30**

TLVWIGIFFC

SEQ ID NO:31

10 SQMMTFFIFL

SEQ ID NO:32

15 VLLLPFVVSC

SEQ ID NO:33

ILCYYWLNTV

20 **SEQ ID NO:34**

TLIVLIITYL

25 **SEQ ID NO:35**

YVLLIRNIFL

SEQ ID NO:36

30 FVFSLVNSFL

SEQ ID NO:37

35 LLLPFVVSCI

SEQ ID NO:38

LIGSVHFFFI

40 **SEQ ID NO:39**

YEMPRHEVYV

45 **SEQ ID NO:40**

LLTRFSAYMV

SEQ ID NO:41

50 IYSGGITKL

SEQ ID NO:42

55 VYYLAEYNL

SEQ ID NO:43

YYWLNTVAL
SEQ ID NO:44

5 IYRLLLMDF
SEQ ID NO:45

10 VYVLLIRNIF
SEQ ID NO:46

CYYWLNTVAL
15 **SEQ ID NO:47**

YYLAEYNLEF
SEQ ID NO:48

20 MLFIMFYSK
SEQ ID NO:49

25 LLIRNIFLK
SEQ ID NO:50

GLQEFDIAR
30 **SEQ ID NO:51**

KMFLIEKLIK
35 **SEQ ID NO:52**

IMLFIMFYSK
SEQ ID NO:53

40 KLIKLDMEK
SEQ ID NO:54

45 VLLIRNIFLK
SEQ ID NO:55

MPRHEVYVL
50 **SEQ ID NO:56**

MPRHEVYVLL
55 **SEQ ID NO:57**

YEMPRHEVY

SEQ ID NO:58

WETLIGQDIY

5 **SEQ ID NO:59**

VHFFFILTL

SEQ ID NO:60

10

LRRIIGMQL

SEQ ID NO:61

15

LLMDFVFSL

SEQ ID NO:62

MQLITSLGL

20

SEQ ID NO:63

FVFSLVNSF

25

SEQ ID NO:64

FILTLIVLI

SEQ ID NO:65

30

WQITEGRKI

SEQ ID NO:66

35

MFYSKNISL

SEQ ID NO:67

FIMFYSKNI

40

SEQ ID NO:68

WRASQMMTF

45

SEQ ID NO:69

FFFILTLIV

SEQ ID NO:70

50

FRLVERYEM

SEQ ID NO:71

55

FFILTLIVL

SEQ ID NO:72

WO 01/23417

PCT/EP00/09500

YAQTLVWI

Claims

1. An isolated polypeptide comprising an amino acid sequence which has at least 70% identity to the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4 over the entire
5 length of of SEQ ID NO:2 or SEQ ID NO:4 respectively.
2. An isolated polypeptide as claimed in claim 1 in which the amino acid sequence has at least 95% identity to SEQ ID NO:2 or SEQ ID NO:4.
- 10 3. The polypeptide as claimed in claim 1 comprising the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4.
4. The isolated polypeptide of SEQ ID NO:2 or SEQ ID NO:4.
- 15 5. A polypeptide comprising an immunogenic fragment of a polypeptide as claimed in any one of claims 1 to 4 in which the immunogenic activity of the immunogenic fragment is substantially the same as the polypeptide of SEQ ID NO:2 or SEQ ID NO:4.
6. A polypeptide as claimed in any of claims 1 to 5 wherein said polypeptide is part of a
20 larger fusion protein.
7. A polypeptide as claimed in any of claims 1 to 6 chemically conjugated to a carrier protein.
- 25 8. An isolated polynucleotide encoding a polypeptide as claimed in any of claims 1 to 6.
9. An isolated polynucleotide comprising a nucleotide sequence encoding a polypeptide that has at least 70% identity to the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4, over the entire length of SEQ ID NO:2 or SEQ ID NO:4 respectively; or a nucleotide
30 sequence complementary to said isolated polynucleotide.
10. An isolated polynucleotide comprising a nucleotide sequence that has at least 70% identity to a nucleotide sequence encoding a polypeptide of SEQ ID NO:2 or SEQ ID NO:4,

over the entire coding region; or a nucleotide sequence complementary to said isolated polynucleotide.

11. An isolated polynucleotide which comprises a nucleotide sequence which has at least
5 70% identity to that of SEQ ID NO:1 or SEQ ID NO:3 over the entire length of SEQ ID
NO:1 or SEQ ID NO:3 respectively; or a nucleotide sequence complementary to said
isolated polynucleotide.

12. The isolated polynucleotide as defined in any one of claims 8 to 11 in which the
10 identity is at least 95%.

13. An isolated polynucleotide selected from:

- (a) a polynucleotide comprising a nucleotide sequence encoding the polypeptide of SEQ
ID NO:2 or SEQ ID NO:4;
- 15 (b) the coding region of the polynucleotide of SEQ ID NO:1 or SEQ ID NO:3; and
- (c) a polynucleotide obtainable by screening an appropriate library under stringent
hybridization conditions with a labeled probe having the sequence of SEQ ID NO:1 or SEQ
ID NO:3 or a fragment thereof said polynucleotide encoding a protein which has similar
immunogenic properties to those of the protein of sequence ID NO:2 or SEQ ID NO:4
20 respectively or a nucleotide sequence complementary to said isolated polynucleotide

14. An expression vector or a recombinant live microorganism comprising an isolated
polynucleotide according to any one of claims 8 - 13.

25 15. A host cell comprising the expression vector of claim 14 or the isolated
polynucleotide of claims 8 to 13.

16. A process for producing a polypeptide of claims 1 to 7 comprising culturing a host
cell of claim 15 under conditions sufficient for the production of said polypeptide and
30 recovering the polypeptide from the culture medium.

17. A vaccine comprising an effective amount of the polypeptide of any one of claims 1
to 7 and a pharmaceutically acceptable carrier.

18. A vaccine comprising an effective amount of the polynucleotide of any one of claims 8 to 13 and a pharmaceutically effective carrier.
- 5 19. A vaccine comprising an effective amount of antigen presenting cells, modified by in vitro loading with a polypeptide of any one of claims 1 to 7, or genetically modified in vitro to express a polypeptide of claims 1 to 7 and a pharmaceutically effective carrier.
20. A vaccine as claimed in any one of claims 17 to 19 which additionally comprises a
10 TH-1 inducing adjuvant.
21. A vaccine as claimed in claim 20 in which the TH-1 inducing adjuvant is selected from the group of adjuvants comprising: 3D-MPL, QS21, a mixture of QS21 and cholesterol, and a CpG oligonucleotide.
- 15 22. An antibody immunospecific for the polypeptide or immunological fragment as claimed in any one of claims 1 to 5.
23. A method for screening to identify compounds which stimulate or which inhibit the
20 function of the polypeptide of any one of claims 1 to 5 which comprises a method selected from the group consisting of:
- (a) measuring the binding of a candidate compound to the said polypeptide (or to the cells or membranes bearing the polypeptide) or a fusion protein thereof by means of a label directly or indirectly associated with the candidate compound;
- 25 (b) measuring the binding of a candidate compound to the said polypeptide (or to the cells or membranes bearing the polypeptide) or a fusion protein thereof in the presence of a labeled competitor;
- (c) testing whether the candidate compound results in a signal generated by activation or inhibition of the said polypeptide, using detection systems appropriate to the cells or cell
30 membranes bearing the polypeptide;
- (d) mixing a candidate compound with a solution containing a polypeptide of any one of claims 1 to 7, to form a mixture, measuring activity of the polypeptide in the mixture, and comparing the activity of the mixture to a standard; or

(e) detecting the effect of a candidate compound on the production of mRNA encoding said polypeptide and said polypeptide in cells, using for instance, an ELISA assay.

24. A method for the treatment of a subject by immunoprophylaxis or therapy
5 comprising *in vitro* induction of immune responses to a molecule of any one of claims 1 to 5, using *in vitro* incubation of the polypeptide of any one of claims 1 to 7 or the polynucleotide of any one of claims 8 to 13 with cells from the immune system of a mammal, and reinfusing these activated immune cells to the mammal for the treatment of disease.

10

25. A method as claimed in claim 24 wherein the treatment is for ovarian or colon cancer.

26. An agonist or antagonist to the polypeptide of claims 1 to 5.

15

27. A compound which is:

(a) an agonist or antagonist to the polypeptide of claims 1 to 5;

(b) isolated polynucleotide of claims 8 to 13; or

(c) a nucleic acid molecule that modulates the expression of the nucleotide sequence

20 encoding the polypeptide of any one of claims 1 to 5;

for use in therapy.

28. A process for diagnosing a disease or a susceptibility to a disease in a subject related to expression or activity of a polypeptide of any one of claims 1 to 5 in a subject

25 comprising analyzing for the presence or amount of said polypeptide in a sample derived from said subject.

29. A process for diagnosing a disease or a susceptibility to a disease in a subject related to expression or activity of a polynucleotide of any one of claims 8 to 13 in a subject

30 comprising analyzing for the presence or amount of said polynucleotide in a sample derived from said subject.

30. A process for diagnosing the presence of colon cancer or a susceptibility to colon cancer in a subject related to expression or activity of a polypeptide of any one of claims 1 to 5 in a subject comprising analyzing for the presence or amount of said polypeptide in a sample derived from said subject.
- 5
31. A process for diagnosing the presence of colon cancer or a susceptibility to colon cancer in a subject related to expression or activity of a polynucleotide of any one of claims 8 to 13 in a subject comprising analyzing for the presence or amount of said polynucleotide in a sample derived from said subject.
- 10
32. An isolated polynucleotide selected from the group consisting of:
- (a) an isolated polynucleotide comprising a nucleotide sequence which has at least 70% identity to SEQ ID NO:5 or SEQ ID NO:7 over the entire length of SEQ ID NO:5 or SEQ ID NO:7 respectively;
 - 15 (b) an isolated polynucleotide comprising the polynucleotide of SEQ ID NO:5 or SEQ ID NO:7;
 - (c) the polynucleotide of SEQ ID NO:5 or SEQ ID NO:7.
33. A live vaccine composition comprising an expression vector or recombinant live
- 20 micro-organism according to claim 14.
34. Use of a polynucleotide as claimed in any one of claims 8 to 13 for the manufacture of a medicament in the treatment of carcinoma.
- 25 35. Use of a polynucleotide as claimed in any one of claims 8 to 13 for the manufacture of a medicament in the treatment of colon carcinoma.
36. Use of a polypeptide as claimed in any one of claims 1 to 7 for the manufacture of a medicament in the treatment of carcinoma.
- 30
37. Use of a polypeptide as claimed in any one of claims 1 to 7 for the manufacture of a medicament in the treatment of colon carcinoma.

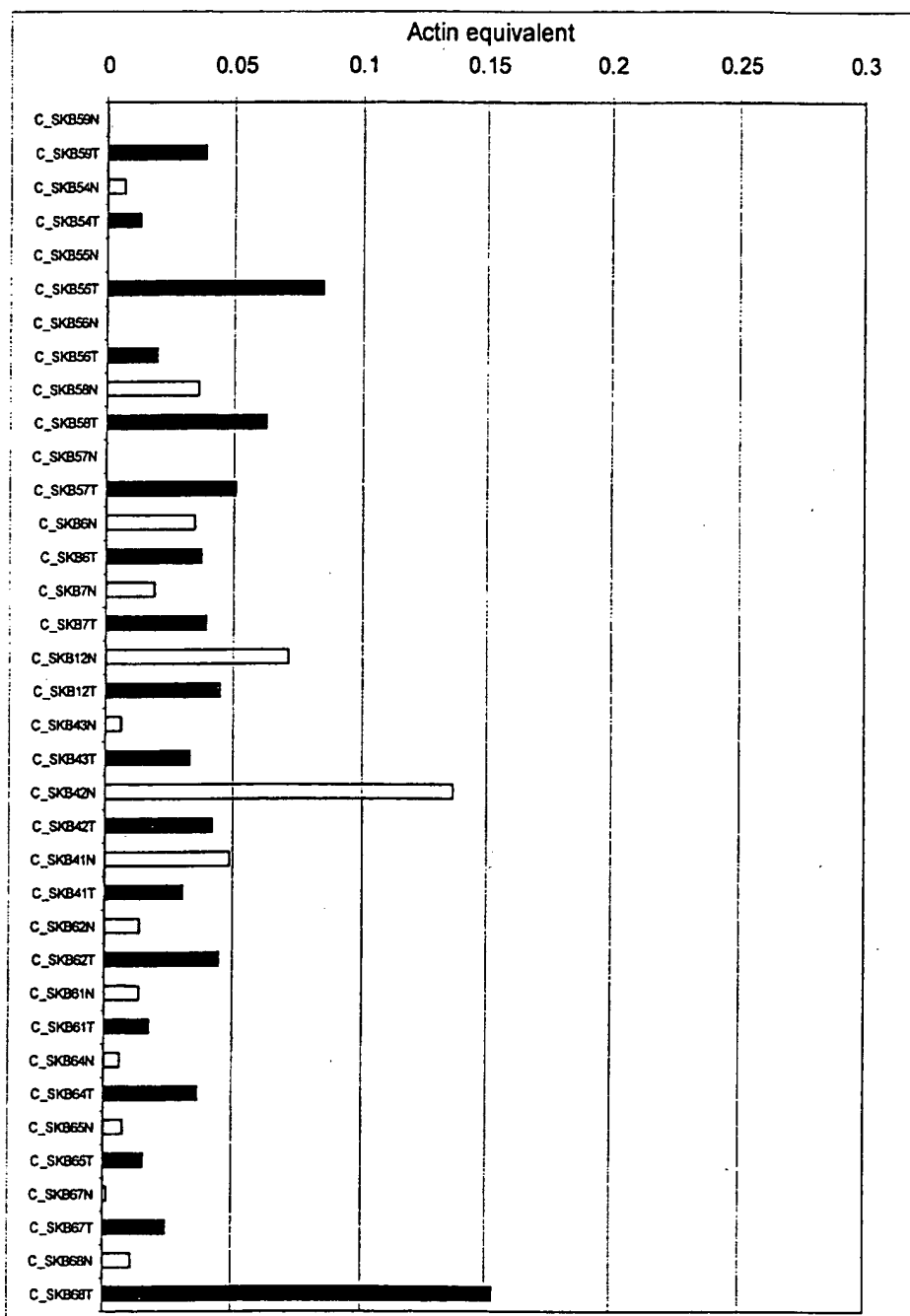
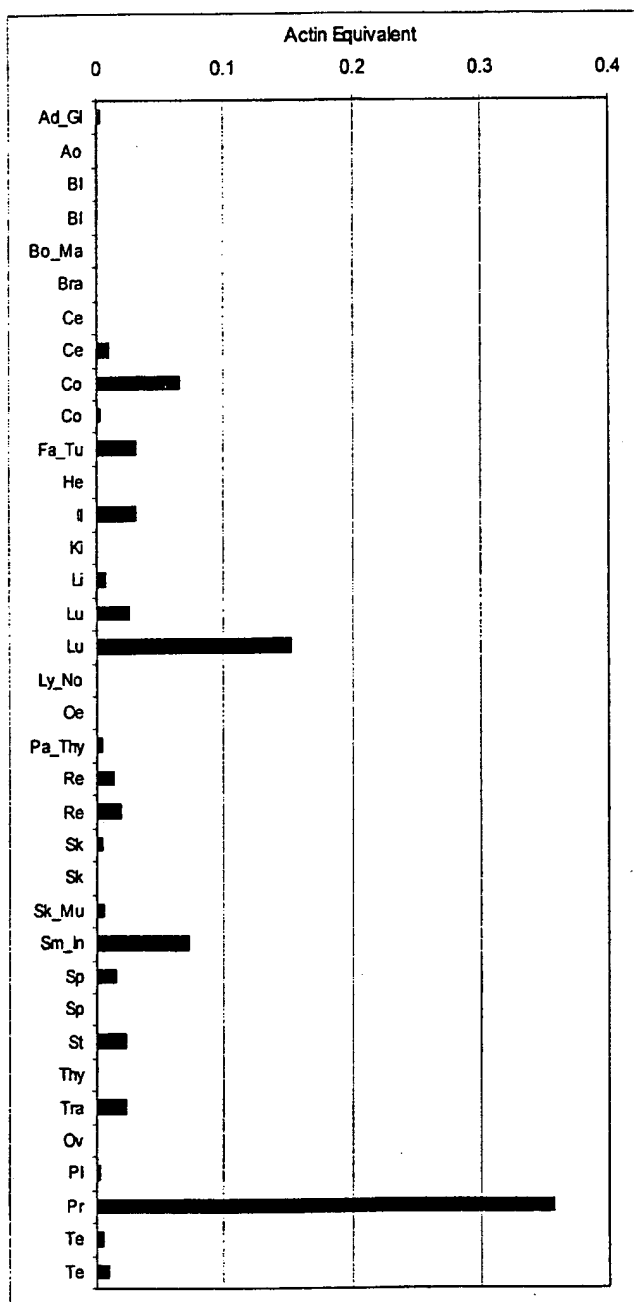


Figure 1: RT PCR data of CASB6411 transcript on colorectal tumours and matched normal colon.

**Legend:**

Ad_Gl: adrenal gland;
 Ao: aorta; Bl: bladder,
 Bo_Ma: bone marrow;
 Bra: brain;
 Ce: cervix; Co: colon;
 Fa_Tu: fallopian tube; He: heart;
 Il: ileon; Ki: kidney; Li: liver;
 Lu: lung; Ly_No: lymph node;
 Oe: oesophagus; Ov: ovary;
 Pa_Thy: parathyroid gland;
 Pl: placenta; Pr: prostate;
 Re: rectum; Sk: skin;
 Sk_Mu: skeletal muscle;
 Sm_In: small intestine;
 Sp: spleen; St: stomach; Te: testis;
 Thy: thyroid gland; Tra: trachea;
 Bre: Breast.

Figure 2: RT PCR data of CASB76411 transcript in a panel of normal tissues.

<110> SmithKline Beecham Biologicals

5

<120> Novel compounds

<130> BC45263

10

<160> 72

<170> FastSEQ for Windows Version 3.0

15

<210> 1

<211> 2407

<212> DNA

<213> Artificial Sequence

20

<400> 1

ttggggaggcca	gaaggcgagac	tgatcacttg	aggccaggag	tttgagacct	catgtctaaa	60
aaaaaaaaaat	tctgtgaggt	gagttttatt	gttattccct	ctctacagat	atggaaactg	120
aggctgagaa	tcgaaccat	tcacaagaca	aaaatcccc	agttggcaga	tccagggttg	180
caagccaggc	ctgtgcagcc	ccaaaaccag	tgcttgttt	accactgtgt	ggtgaccaca	240
ccgctccagg	ccaacagctt	ggggctaagt	cttccagttg	ccctttacca	ttaaataata	300
gggctgcctt	ttgttgaagc	cctgcactcc	cagtgacggc	cataataaac	ttcaggtgtt	360
ctgctttctg	ccttctctag	catggccaag	tatttccgga	acaacttcat	taatccccac	420
atttactccg	gagggatcac	caagctgata	ttttgctggg	acttcaactg	cactcatgaa	480
aaagctgtga	agctaaaaca	gaagaatctt	agcactgata	taagggagaa	cctgtcagag	540
ctccgtcagg	agaattccaa	gttgacgttc	aatcagctgc	tgacccgctt	ctctgcctac	600
atggtagcct	gggttgtctc	tacaggagtg	gccatagcct	gctgtgcagc	cgtttattac	660
ctggctgagt	acaacttaga	gttcctgaag	acacacagta	accctggggc	ggtgtctgta	720
ctgcctttcg	ttgtgtcctg	cattaatctg	gccgtgccat	gcacttactc	catgtctcag	780
cttctggaga	ggtacgagat	gccacggcac	gaagttacg	ttctctctgt	ccgaaacatc	840
tttttgaaaa	tatcaatcat	tggcattctt	tgttactatt	ggctcaacac	cgtaggcctg	900
tctggtgaag	agtgttgga	aacctcatt	ggccaggaca	tctaccggct	ccttctgtat	960
gatttttgtt	tctctttaag	caattccttc	ctgggggagt	tcttgaggag	aatcattggg	1020
atgcaactga	tcacaagctt	tgccctcag	gagtttgaca	ttgccaggaa	cgttctagaa	1080
ctctacttat	cacaaactct	ggtgtggatt	ggcatcttct	tctgccccct	gctgcccttt	1140
atccaaatga	ttatgctttt	catcatgttc	tactccaaaa	atatcagcct	gatgatgaat	1200
ttccagcctc	cgagcaaaag	ctggcggggc	tcacagatga	tgactttctt	catcttcttg	1260
ctctttttcc	catccttcac	cggggtcttg	ggcaccttg	ccatcaccat	ctggagattg	1320
aagcctttac	ctcagcttgg	ccttttctga	tgctgtccct	tcttcattca	ctccactctc	1380
agctggatcg	acaccctaag	tacacggcct	ggctacctgt	gggttgtttg	gatctatcgg	1440
aacctcattg	gaagtgtgca	cttctttttc	atcctcaccc	tcattgtgct	aatcatcacc	1500
tatctttaact	ggcagatcac	agaggggaag	aagattatga	taaggctgtg	cctgagcag	1560
atcatttaagt	agggcaaaag	taaaatgttc	gatgatgaaa	aattgtacaa	gctgcaggat	1620
atggagaaga	aagcaaaacc	cagctcactt	gttcttgaaa	ggagagaggt	ggagcaacaa	1680
ggctttttgc	atttggggga	acatgatggc	agtcttgact	tgcgatctag	aagatcagtt	1740
caagaaggta	atccaagggc	ctgatgactc	ttttggtaac	cagacaccaa	tcaataaagg	1800
gagggagacg	aaaatggaat	gatttctctc	atgcccactg	tgccctttag	aactgcccg	1860
aagaaataacc	aaggcttttag	caggagcgg	aaactgacta	ccatgtaatt	atcaaagtaa	1920
aattgggcat	tccatgctat	ttttaatacc	tggttggctg	atttttcaag	acaaaaatac	1980
tggggttttc	caataaagat	tgttgttaata	ttgaaatgag	cctacaaaaa	cctaggaaga	2040
gataactagg	gaataatgta	tattattctc	aggaaatgtg	tgagggaatg	atttgttctt	2100
agaaatctct	cctgccagac	tcccagacc	tggaaaaggt	ttagaactctg	ttgtctaaga	2160
aagtggtcca	tctgaataaa	acatgtaata	ctccagcagg	gatatgaagc	ctctgaattg	2220
tagaacctgc	atttatttgt	gactttgaac	taaagacatc	ccccatgtcc	caaaggtgga	2280
atacaccagg	aggtctcatt	tctgaacttt	cttgcgctact	gattacatga	gtctttggag	2340
tcggggatgg	aggaggttct	gcccctgtga	ggtgttatac	atgaccatca	aagtcctacg	2400
tcaagct						2407

<210> 2

<211> 460

65

<212> PRT

<213> Artificial Sequence

<400> 2
 Met Ala Lys Tyr Phe Arg Asn Asn Phe Ile Asn Pro His Ile Tyr Ser
 1 5 10
 5 Gly Gly Ile Thr Lys Leu Ile Phe Cys Trp Asp Phe Thr Val Thr His
 20 25 30
 Glu Lys Ala Val Lys Leu Lys Gln Lys Asn Leu Ser Thr Glu Ile Arg
 35 40 45
 10 Glu Asn Leu Ser Glu Leu Arg Gln Glu Asn Ser Lys Leu Thr Phe Asn
 50 55 60
 Gln Leu Leu Thr Arg Phe Ser Ala Tyr Met Val Ala Trp Val Val Ser
 65 70 75 80
 Thr Gly Val Ala Ile Ala Cys Cys Ala Ala Val Tyr Tyr Leu Ala Glu
 85 90 95
 15 Tyr Asn Leu Glu Phe Leu Lys Thr His Ser Asn Pro Gly Ala Val Leu
 100 105 110
 Leu Leu Pro Phe Val Val Ser Cys Ile Asn Leu Ala Val Pro Cys Ile
 115 120 125
 20 Tyr Ser Met Phe Arg Leu Val Glu Arg Tyr Glu Met Pro Arg His Glu
 130 135 140
 Val Tyr Val Leu Leu Ile Arg Asn Ile Phe Leu Lys Ile Ser Ile Ile
 145 150 155 160
 Gly Ile Leu Cys Tyr Tyr Trp Leu Asn Thr Val Ala Leu Ser Gly Glu
 165 170 175
 25 Glu Cys Trp Glu Thr Leu Ile Gly Gln Asp Ile Tyr Arg Leu Leu Leu
 180 185 190
 Met Asp Phe Val Phe Ser Leu Val Asn Ser Phe Leu Gly Glu Phe Leu
 195 200 205
 Arg Arg Ile Ile Gly Met Gln Leu Ile Thr Ser Leu Gly Leu Gln Glu
 210 215 220
 30 Phe Asp Ile Ala Arg Asn Val Leu Glu Leu Ile Tyr Ala Gln Thr Leu
 225 230 235 240
 Val Trp Ile Gly Ile Phe Phe Cys Pro Leu Leu Pro Phe Ile Gln Met
 245 250 255
 35 Ile Met Leu Phe Ile Met Phe Tyr Ser Lys Asn Ile Ser Leu Met Met
 260 265 270
 Asn Phe Gln Pro Pro Ser Lys Ala Trp Arg Ala Ser Gln Met Met Thr
 275 280 285
 40 Phe Phe Ile Phe Leu Leu Phe Phe Pro Ser Phe Thr Gly Val Leu Cys
 290 295 300
 Thr Leu Ala Ile Thr Ile Trp Arg Leu Lys Pro Ser Ala Asp Cys Gly
 305 310 315 320
 Pro Phe Arg Gly Leu Pro Leu Phe Ile His Ser Ile Tyr Ser Trp Ile
 325 330 335
 45 Asp Thr Leu Ser Thr Arg Pro Gly Tyr Leu Trp Val Val Trp Ile Tyr
 340 345 350
 Arg Asn Leu Ile Gly Ser Val His Phe Phe Phe Ile Leu Thr Leu Ile
 355 360 365
 50 Val Leu Ile Ile Thr Tyr Leu Tyr Trp Gln Ile Thr Glu Gly Arg Lys
 370 375 380
 Ile Met Ile Arg Leu Leu His Glu Gln Ile Ile Asn Glu Gly Lys Asp
 385 390 395 400
 Lys Met Phe Leu Ile Glu Lys Leu Ile Lys Leu Gln Asp Met Glu Lys
 405 410 415
 55 Lys Ala Asn Pro Ser Ser Leu Val Leu Glu Arg Arg Glu Val Glu Gln
 420 425 430
 Gln Gly Phe Leu His Leu Gly Glu His Asp Gly Ser Leu Asp Leu Arg
 435 440 445
 60 Ser Arg Arg Ser Val Gln Glu Gly Asn Pro Arg Ala
 450 455 460

<210> 3
 <211> 2521
 <212> DNA
 65 <213> Artificial Sequence

<400> 3

	tggggaggca	gaaggcagac	tgatcacttg	aggccaggag	tttgagacct	catgtctaaa	60
	aaaaaaaaat	tctgtgaggt	gagttttatt	gttattccct	ctctacagat	atggaaaactg	120
	aggctgagaa	tcagaacccat	tcacaagaca	aaaatccccc	agttggcaga	tccagggttg	180
5	caagccaggc	ctgtgcagcc	ccaaaaccag	tgcttgttta	accactgtgt	ggtgaccaca	240
	ccgctccagg	ccaacagctt	ggggctaagt	cttcacgttg	cctttcacca	ttaaataata	300
	gggctgccct	ttgttgaagc	cctgcactcc	cagtgcgggc	cataataacc	ttcagggtgtt	360
	ctgctttctg	ccttctctag	catggccaag	tatttccgga	acaacttcat	taatccccac	420
	atttactccg	gagggatcac	caagctgatc	ttttgctggg	acttcaactgt	cactcatgaa	480
10	aaagctgtga	agctaaaaca	gaagaatctt	agcactgaga	taaggagaga	cctgtcagag	540
	ctccgtcagg	agaattccaa	gttgacgttc	aatcagctgc	tgaccgctt	ctctgcctac	600
	atggtagcct	gggttgtctc	tacaggagtg	gccatagcct	gctgtgcagc	cgtttattac	660
	ctggctgagt	acaacttaga	gttcctgaag	acacacagta	accctggggc	ggtgctgtta	720
	ctgccttttg	ttgtgtcctg	cattctggcc	gtgccatgca	tctactccat	gttcaggctt	780
15	gtggagaggt	acgagatgcc	acggcacgaa	gtctacgttc	tcctgatccg	caggggattg	840
	atgtagtctc	caagtatggg	atgtacagat	gggcaggcag	tgcacgcaca	aaggctcctg	900
	ggctgaggag	gggactgaaa	tcattccagcg	ttcccccttag	tcaagctaaa	catctttttg	960
	aaaatatcaa	actttggcat	tctttgttac	tattggctca	acaccgtggc	cctgtctggt	1020
	gaagagtgtt	gggaaacccct	cattggccag	gacatctacc	ggctccttct	gatggatttt	1080
20	gtgttctctt	tagtcaattc	cttcctgggg	gagtttctga	ggagaatcat	tgggatgcaa	1140
	ctgatcacia	gtcctggcct	tcaggagttt	gacattgccca	ggaacgttct	agaactgatc	1200
	tatgcacaaa	ctctggtgtg	gattggcatc	ttcttctgcc	ccctgctgcc	ctttatccaa	1260
	atgattatgc	ttttcatcat	gttctactcc	aaaaatatca	gcctgatgat	gaatttccag	1320
	cctccgagca	aagcctggcg	ggcctcacag	atgatgactt	tcttcatctt	cttgctcttt	1380
25	ttcccatcct	tcaccggggt	cttgtgcacc	ctggccatca	ccatctggag	attgaagcct	1440
	tcagctgact	gtggcccttt	tcgaggctct	cctctcttca	ttcactccat	ctacagctgg	1500
	atcgacacc	taagtaacag	gcctggctac	ctgtgggttg	tttggatcta	tcggaacctc	1560
	attggaagt	tgcacttctt	tttcatcctc	accctcattg	tgctaatacat	cacctatctt	1620
	tactggcaga	tcacagaggg	aagggaagatt	atgataaggc	tgctccatga	gcagatcatt	1680
30	aatgagggca	aagataaaat	gttcctgata	gaaaaattga	tcaagctgca	ggatatggag	1740
	aagaaagcaa	accccagctc	acttgttctg	gaaaggagag	aggtggagca	acaaggcttt	1800
	ttgcatttgg	gggaacatga	tggcagtctt	gacttgcgat	ctagaagatc	agttcaagaa	1860
	ggtaaatcaa	gggcctgatg	actcttttgg	taaccagaca	ccaatcaaat	aaggggagga	1920
	gacgaaaaatg	gaatgatttc	ttccatgcca	cctgtgcctt	taggaactgc	ccagaagaaa	1980
35	atccaaggct	ttagccagga	gcggaactg	actaccatgt	aattatcaaa	gtaaaattgg	2040
	gcattccatg	ctatttttaa	tacctggatt	gctgattttt	caagacaaaa	tacttggggg	2100
	tttccaataa	agattgttgt	aatattgaaa	tgagcctaca	aaaacctagg	aagagataac	2160
	tagggaataa	tgtatattat	cttcaagaaa	tgtgtgcagg	aatgatttgt	tcttagaaaat	2220
	ctctccttgc	agacttccca	gacctggcaa	aggttttagaa	actgttgcta	agaaaagtgg	2280
40	tccatcctga	ataaacatgt	aatactccag	cagggatatg	aagcctctga	attgtagaac	2340
	ctgcatttat	ttgtgacttt	gaactaaaga	catcccccat	gtcccaaaag	tggaaatacaa	2400
	ccagaggtct	catctctgaa	ctttcttgcg	tactgattac	atgagtcttt	ggagtcgggg	2460
	atggaggagg	ttctgcccct	gtgagggtgt	atacatgacc	atcaaagtcc	tacgtcaagc	2520
45	t						2521

<210> 4
 <211> 154
 <212> PRT
 <213> Artificial Sequence

50 <400> 4

	Met	Ala	Lys	Tyr	Phe	Arg	Asn	Asn	Phe	Ile	Asn	Pro	His	Ile	Tyr	Ser
	1				5				10					15		
	Gly	Gly	Ile	Thr	Lys	Leu	Ile	Phe	Cys	Trp	Asp	Phe	Thr	Val	Thr	His
55				20					25					30		
	Glu	Lys	Ala	Val	Lys	Leu	Lys	Gln	Lys	Asn	Leu	Ser	Thr	Glu	Ile	Arg
			35					40					45			
	Glu	Asn	Leu	Ser	Glu	Leu	Arg	Gln	Glu	Asn	Ser	Lys	Leu	Thr	Phe	Asn
		50				55						60				
60	Gln	Leu	Leu	Thr	Arg	Phe	Ser	Ala	Tyr	Met	Val	Ala	Trp	Val	Val	Ser
	65				70				75					80		
	Thr	Gly	Val	Ala	Ile	Ala	Cys	Cys	Ala	Ala	Val	Tyr	Tyr	Leu	Ala	Glu
				85					90					95		
	Tyr	Asn	Leu	Glu	Phe	Leu	Lys	Thr	His	Ser	Asn	Pro	Gly	Ala	Val	Leu
65			100					105					110			
	Leu	Leu	Pro	Phe	Val	Val	Ser	Cys	Ile	Leu	Ala	Val	Pro	Cys	Ile	Tyr

	115	120	125
	Ser Met Phe Arg Leu Val Glu Arg Tyr Glu Met	Pro Arg His Glu Val	
	130	135	140
5	Tyr Val Leu Leu Ile Arg Arg Gly Leu Met		
	145	150	
	<210> 5		
	<211> 1960		
	<212> DNA		
10	<213> Artificial Sequence		
	<400> 5		
	atcttttgcg gggacttcac tgtcactcat gaaaaagctg tgaagctaaa acagaagaat	60	
	cttagcactg agataaggga gaacctgtca gagctccgtc aggagaattc caagttgacg	120	
15	ttcaatcagc tgctgaccgc cttctctgcc tacatggtag cctgggttgt ctctacagga	180	
	gtggccatag cctgctgtgc agccgtttat tacctggctg agtacaactt agagttcctg	240	
	aagacacaca gtaacctcgg ggcggtgctg ttactgcctt tcggtgtgtc ctgcattaat	300	
	ctggccgtgc catgcactca ctccatgttc aggccttggtg agaggtaacg gatgccacgg	360	
	cacgaagtct acgttctcct gatccgaaac atctttttga aaatatcaat cattggcatt	420	
20	ctttgttact attggctcaa caccgtggcc ctgtctggtg aagagtgttg ggaaccctc	480	
	attggccagg acatctaccg gctccttctg atggattttg tgttctcttt agtcaattcc	540	
	ttcctggggg agtttctgag gagaatcatt gggatgcaac tgatcacaag tcttggcctt	600	
	caggagtgtg acattgccag gaacgttcta gaactgatct atgcacaaac tctgggtgtg	660	
	attggcatct tcttctgccc cctgctgccc tttatccaaa tgattatgct tttcatcatg	720	
25	ttctactcca aaaatatcag cctgatgatg aatttccagc ctccgagcaa agcctggcgg	780	
	gcctcacaga tgatgacttt cttcatcttc ttgctctttt tcccatcttt caccggggtc	840	
	ttgtgcaccc tggccatcac catctggaga ttgaagcctt cagctgactg tggccctttt	900	
	cgaggctctg ctctcttcat tcaactccatc tacagctgga tcgacaccct aagtacacgg	960	
	cctggctacc tgtgggttgt ttggatctat cggaaacctca ttggaagtgt gcacttcttt	1020	
30	ttcatcctca cctcatttgt gctgatcatt acctatcttt actggcagat cacagaggga	1080	
	aggaaagatta tgataaggct gctccatgag cagatcatta atgagggcaa agataaaatg	1140	
	ttcctgatag aaaaattgat caagctgcag gatattggaga agaaagcaaa cccagctca	1200	
	cttgttctgg aaaggagaga ggtggagcaa caaggctttt tgcatttggg ggaacatgat	1260	
	ggcagctctg acttgcgac tagaagatca gttcaagaag gtaatccaag ggctgatga	1320	
35	ctcttttggg aaccagacac caatcaaaata aggggaggag atgaaaatgg aatgatttct	1380	
	tccatgccac ctgtgccttt aggaactgcc cagaagaaaa tccaaggctt tagccaggag	1440	
	cgaaaactga ctaccatgta attatcaaaag taaaattggg cattccatgc tatttttaat	1500	
	acctggattg ctgatttttc aagacaaaat acttgggggt ttccaataaa gattgttgta	1560	
	atattgaaat ggcctacaa aaacctagga agagataact agggaataat gtatattatc	1620	
40	ttcaagaaat gtgtgcagga atgattggtt cttagaaatc tctcctgccg gacttcccag	1680	
	acctggcaaa ggtttagaaa ctgttgctaa gaaaagtggg ccactctgaa taacatgta	1740	
	atactccagc agggatagta agcctctgaa ttgtagaacc tgcatttatt tgtgactttg	1800	
	aactaaagac atccccatg tcccaaagggt ggaatacaac cagaggtctc atctctgaac	1860	
	tttcttgcgt actgattaca tgagtctttg gagtctggga tggaggaggt tctgccctg	1920	
45	tgagggtgta tacatgacca tcaaagtcct acgtcaagct	1960	
	<210> 6		
	<211> 438		
	<212> PRT		
50	<213> artificial sequence		
	<400> 6		
	Ile Phe Cys Trp Asp Phe Thr Val Thr His Glu Lys Ala Val Lys Leu		
	1 5 10 15		
55	Lys Gln Lys Asn Leu Ser Thr Glu Ile Arg Glu Asn Leu Ser Glu Leu		
	20 25 30		
	Arg Gln Glu Asn Ser Lys Leu Thr Phe Asn Gln Leu Leu Thr Arg Phe		
	35 40 45		
60	Ser Ala Tyr Met Val Ala Trp Val Val Ser Thr Gly Val Ala Ile Ala		
	50 55 60		
	Cys Cys Ala Ala Val Tyr Tyr Leu Ala Glu Tyr Asn Leu Glu Phe Leu		
	65 70 75 80		
	Lys Thr His Ser Asn Pro Gly Ala Val Leu Leu Leu Pro Phe Val Val		
	85 90 95		
65	Ser Cys Ile Asn Leu Ala Val Pro Cys Ile Tyr Ser Met Phe Arg Leu		
	100 105 110		

	Val	Glu	Arg	Tyr	Glu	Met	Pro	Arg	His	Glu	Val	Tyr	Val	Leu	Leu	Ile	
		115						120					125				
	Arg	Asn	Ile	Phe	Leu	Lys	Ile	Ser	Ile	Ile	Gly	Ile	Leu	Cys	Tyr	Tyr	
		130					135					140					
5	Trp	Leu	Asn	Thr	Val	Ala	Leu	Ser	Gly	Glu	Glu	Cys	Trp	Glu	Thr	Leu	
		145				150					155					160	
	Ile	Gly	Gln	Asp	Ile	Tyr	Arg	Leu	Leu	Leu	Met	Asp	Phe	Val	Phe	Ser	
				165						170					175		
	Leu	Val	Asn	Ser	Phe	Leu	Gly	Glu	Phe	Leu	Arg	Arg	Ile	Ile	Gly	Met	
10			180						185					190			
	Gln	Leu	Ile	Thr	Ser	Leu	Gly	Leu	Gln	Glu	Phe	Asp	Ile	Ala	Arg	Asn	
			195					200					205				
	Val	Leu	Glu	Leu	Ile	Tyr	Ala	Gln	Thr	Leu	Val	Trp	Ile	Gly	Ile	Phe	
		210					215					220					
15	Phe	Cys	Pro	Leu	Leu	Pro	Phe	Ile	Gln	Met	Ile	Met	Leu	Phe	Ile	Met	
						230					235					240	
	Phe	Tyr	Ser	Lys	Asn	Ile	Ser	Leu	Met	Met	Asn	Phe	Gln	Pro	Pro	Ser	
					245					250					255		
	Lys	Ala	Trp	Arg	Ala	Ser	Gln	Met	Met	Thr	Phe	Phe	Ile	Phe	Leu	Leu	
20				260					265					270			
	Phe	Phe	Pro	Ser	Phe	Thr	Gly	Val	Leu	Cys	Thr	Leu	Ala	Ile	Thr	Ile	
				275				280					285				
	Trp	Arg	Leu	Lys	Pro	Ser	Ala	Asp	Cys	Gly	Pro	Phe	Arg	Gly	Leu	Pro	
		290					295					300					
25	Leu	Phe	Ile	His	Ser	Ile	Tyr	Ser	Trp	Ile	Asp	Thr	Leu	Ser	Thr	Arg	
						310					315					320	
	Pro	Gly	Tyr	Leu	Trp	Val	Val	Trp	Ile	Tyr	Arg	Asn	Leu	Ile	Gly	Ser	
						325					330				335		
	Val	His	Phe	Phe	Phe	Ile	Leu	Thr	Leu	Ile	Val	Leu	Ile	Ile	Thr	Tyr	
30				340					345					350			
	Leu	Tyr	Trp	Gln	Ile	Thr	Glu	Gly	Arg	Lys	Ile	Met	Ile	Arg	Leu	Leu	
				355				360					365				
	His	Glu	Gln	Ile	Ile	Asn	Glu	Gly	Lys	Asp	Lys	Met	Phe	Leu	Ile	Glu	
				370			375					380					
35	Lys	Leu	Ile	Lys	Leu	Gln	Asp	Met	Glu	Lys	Lys	Ala	Asn	Pro	Ser	Ser	
						390					395					400	
	Leu	Val	Leu	Glu	Arg	Arg	Glu	Val	Glu	Gln	Gln	Gly	Phe	Leu	His	Leu	
					405					410					415		
	Gly	Glu	His	Asp	Gly	Ser	Leu	Asp	Leu	Arg	Ser	Arg	Arg	Ser	Val	Gln	
40				420					425					430			
	Glu	Gly	Asn	Pro	Arg	Ala											
				435													
	<210> 7																
45	<211> 1219																
	<212> DNA																
	<213> Artificial Sequence																
	<400> 7																
50	ctgatgatga	atttccagcc	tccgagcaaa	gcctggcggg	cctcacagat	gatgactttc											60
	ttcatcttct	tgctcttttt	cccatctttc	accggggtct	tgtgcaccct	ggccatcacc											120
	atctggagat	tgaagccttc	agctgactgt	ggcccttttc	gaggtctgcc	tctcttcatt											180
	cactccatct	acagctggat	cgacacccta	agtacacggc	ctggctacct	gtgggttggt											240
	tggatctatc	ggaacctcat	tggaagtgtg	cacttctttt	tcatectcac	cctcattgtg											300
55	ctgatcatca	cctatcttta	ctggcagatc	acagagggaa	ggaagattat	gataaggctg											360
	ctccatgagc	agatcattaa	tgaggggcaaa	gataaaatgt	tcctgataga	aaaattgatc											420
	aagctgcagg	atatggagaa	gaaagcaaac	cccagctcac	ttgttctgga	aaggagagag											480
	gtggagcaac	aaggcttttt	gcatttgggg	gaacatgatg	gcagtcctga	cttgcgatct											540
	agaagatcag	ttcaagaagg	taatccaagg	gcctgatgac	tcttttggtg	accagacacc											600
60	aatcaataaa	ggggaggaga	tgaaaatgga	atgatttctt	ccatgccacc	tgtgccttta											660
	ggaactgccc	agaagaaaat	caaaggcttt	agccaggagc	ggaaactgac	taccatgtaa											720
	ttatcaaaagt	aaaattgggc	attccatgct	atttttaata	cctggattgc	tgatttttca											780
	agacaaaaata	cttgggggttt	tccaataaaag	attgttgtaa	tattgaaatg	agcctacaaa											840
	aacctaggaa	gagataaacta	gggaataatg	tatattatct	tcaagaaaatg	tgtgcaggaa											900
65	tgattggttc	ttagaaaatct	ctcctgccag	acttcccaga	cctggcaaag	gtttagaaac											960
	tgttgctaag	aaaagtggct	catcctgaat	aaacatgtaa	tactccagca	gggatatgaa											1020

gcctctgaat tgtagaacct gcatttattt gtgactttga actaaagaca tcccccatgt 1080
 cccaaagggtg gaatacaacc agagggtctca tctctgaact ttcttgcgta ctgattacat 1140
 gagtctttgg agtcggggat ggaggaggtt ctgcccctgt gaggtgttat acatgaccat 1200
 caaagtcccta cgtaagct 1219

5
 <210> 8
 <211> 191
 <212> PRT
 <213> Artificial Sequence

10
 <400> 8
 Leu Met Met Asn Phe Gln Pro Pro Ser Lys Ala Trp Arg Ala Ser Gln
 1 5 10 15
 Met Met Thr Phe Phe Ile Phe Leu Leu Phe Phe Pro Ser Phe Thr Gly
 15 20 25 30
 Val Leu Cys Thr Leu Ala Ile Thr Ile Trp Arg Leu Lys Pro Ser Ala
 35 40 45
 Asp Cys Gly Pro Phe Arg Gly Leu Pro Leu Phe Ile His Ser Ile Tyr
 50 55 60
 20 Ser Trp Ile Asp Thr Leu Ser Thr Arg Pro Gly Tyr Leu Trp Val Val
 65 70 75 80
 Trp Ile Tyr Arg Asn Leu Ile Gly Ser Val His Phe Phe Phe Ile Leu
 85 90 95
 Thr Leu Ile Val Leu Ile Ile Thr Tyr Leu Tyr Trp Gln Ile Thr Glu
 100 105 110
 25 Gly Arg Lys Ile Met Ile Arg Leu Leu His Glu Gln Ile Ile Asn Glu
 115 120 125
 Gly Lys Asp Lys Met Phe Leu Ile Glu Lys Leu Ile Lys Leu Gln Asp
 130 135 140
 30 Met Glu Lys Lys Ala Asn Pro Ser Ser Leu Val Leu Glu Arg Arg Glu
 145 150 155 160
 Val Glu Gln Gln Gly Phe Leu His Leu Gly Glu His Asp Gly Ser Leu
 165 170 175
 35 Asp Leu Arg Ser Arg Arg Ser Val Gln Glu Gly Asn Pro Arg Ala
 180 185 190

40
 <210> 9
 <211> 10
 <212> PRT
 <213> Artificial Sequence

45
 <400> 9
 Ile Thr Glu Gly Arg Lys Ile Met Ile Arg
 1 5 10

50
 <210> 10
 <211> 9
 <212> PRT
 <213> Artificial Sequence

55
 <400> 10
 Leu Leu Met Asp Phe Val Phe Ser Leu
 1 5

60
 <210> 11
 <211> 9
 <212> PRT
 <213> Artificial Sequence

65
 <400> 11
 Phe Leu Leu Phe Phe Pro Ser Phe Thr
 1 5

70
 <210> 12
 <211> 9
 <212> PRT

```

      <213> Artificial Sequence
      <400> 12
5      Gln Met Met Thr Phe Phe Ile Phe Leu
      1              5

      <210> 13
      <211> 9
      <212> PRT
10     <213> Artificial Sequence

      <400> 13
      Met Met Thr Phe Phe Ile Phe Leu Leu
      1              5
15

      <210> 14
      <211> 9
      <212> PRT
      <213> Artificial Sequence
20

      <400> 14
      Phe Leu Ile Glu Lys Leu Ile Lys Leu
      1              5
25

      <210> 15
      <211> 9
      <212> PRT
      <213> Artificial Sequence
30

      <400> 15
      Val Leu Leu Ile Arg Asn Ile Phe Leu
      1              5
35

      <210> 16
      <211> 9
      <212> PRT
      <213> Artificial Sequence

      <400> 16
40     Leu Val Trp Ile Gly Ile Phe Phe Cys
      1              5

      <210> 17
      <211> 9
      <212> PRT
45     <213> Artificial Sequence

      <400> 17
      Thr Leu Ala Ile Thr Ile Trp Arg Leu
50     1              5

      <210> 18
      <211> 9
      <212> PRT
55     <213> Artificial Sequence

      <400> 18
      Leu Ile Phe Cys Trp Asp Phe Thr Val
      1              5
60

      <210> 19
      <211> 9
      <212> PRT
      <213> Artificial Sequence
65

      <400> 19

```

```

Phe Leu Gly Glu Phe Leu Arg Arg Ile
1           5

    <210> 20
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 20
10  Leu Leu Leu Pro Phe Val Val Ser Cys
    1           5

    <210> 21
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 21
20  Leu Leu Thr Arg Phe Ser Ala Tyr Met
    1           5

    <210> 22
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 22
30  Lys Leu Ile Phe Cys Trp Asp Phe Thr
    1           5

    <210> 23
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 23
35  Leu Leu Leu Met Asp Phe Val Phe Ser Leu
    1           5           10

    <210> 24
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 24
45  Leu Leu Met Asp Phe Val Phe Ser Leu Val
    1           5           10

    <210> 25
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 25
55  Gln Met Met Thr Phe Phe Ile Phe Leu Leu
    1           5           10

    <210> 26
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 26
65  Tyr Leu Ala Glu Tyr Asn Leu Glu Phe Leu
    1           5           10

```


<210> 27
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 5
 <400> 27
 Lys Leu Ile Phe Cys Trp Asp Phe Thr Val
 1 5 10
 10
 <210> 28
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 15
 <400> 28
 Leu Leu Phe Phe Pro Ser Phe Thr Gly Val
 1 5 10
 20
 <210> 29
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 25
 <400> 29
 Gln Leu Leu Thr Arg Phe Ser Ala Tyr Met
 1 5 10
 30
 <210> 30
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 35
 <400> 30
 Thr Leu Val Trp Ile Gly Ile Phe Phe Cys
 1 5 10
 40
 <210> 31
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 45
 <400> 31
 Ser Gln Met Met Thr Phe Phe Ile Phe Leu
 1 5 10
 50
 <210> 32
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 55
 <400> 32
 Val Leu Leu Leu Pro Phe Val Val Ser Cys
 1 5 10
 60
 <210> 33
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 65
 <400> 33
 Ile Leu Cys Tyr Tyr Trp Leu Asn Thr Val
 1 5 10
 <210> 34
 <211> 10
 <212> PRT

```

      <213> Artificial Sequence
      <400> 34
5    Thr Leu Ile Val Leu Ile Ile Thr Tyr Leu
      1             5             10

      <210> 35
      <211> 10
      <212> PRT
10    <213> Artificial Sequence

      <400> 35
      Tyr Val Leu Leu Ile Arg Asn Ile Phe Leu
      1             5             10
15

      <210> 36
      <211> 10
      <212> PRT
      <213> Artificial Sequence
20

      <400> 36
      Phe Val Phe Ser Leu Val Asn Ser Phe Leu
      1             5             10
25

      <210> 37
      <211> 10
      <212> PRT
      <213> Artificial Sequence
30

      <400> 37
      Leu Leu Leu Pro Phe Val Val Ser Cys Ile
      1             5             10
35

      <210> 38
      <211> 10
      <212> PRT
      <213> Artificial Sequence

      <400> 38
40    Leu Ile Gly Ser Val His Phe Phe Phe Ile
      1             5             10

      <210> 39
      <211> 10
      <212> PRT
45    <213> Artificial Sequence

      <400> 39
      Tyr Glu Met Pro Arg His Glu Val Tyr Val
50    1             5             10

      <210> 40
      <211> 10
      <212> PRT
55    <213> Artificial Sequence

      <400> 40
      Leu Leu Thr Arg Phe Ser Ala Tyr Met Val
      1             5             10
60

      <210> 41
      <211> 9
      <212> PRT
      <213> Artificial Sequence
65

      <400> 41

```

```

Ile Tyr Ser Gly Gly Ile Thr Lys Leu
1           5

    <210> 42
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 42
10 Val Tyr Tyr Leu Ala Glu Tyr Asn Leu
   1           5

    <210> 43
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 43
20 Tyr Tyr Trp Leu Asn Thr Val Ala Leu
   1           5

    <210> 44
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 44
30 Ile Tyr Arg Leu Leu Leu Met Asp Phe
   1           5

    <210> 45
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 45
35 Val Tyr Val Leu Leu Ile Arg Asn Ile Phe
   1           5           10

    <210> 46
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 46
45 Cys Tyr Tyr Trp Leu Asn Thr Val Ala Leu
   1           5           10

    <210> 47
    <211> 10
    <212> PRT
    <213> Artificial Sequence

    <400> 47
55 Tyr Tyr Leu Ala Glu Tyr Asn Leu Glu Phe
   1           5           10

    <210> 48
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 48
65 Met Leu Phe Ile Met Phe Tyr Ser Lys
   1           5

```

<210> 49
 <211> 9
 <212> PRT
 <213> Artificial Sequence
 5
 <400> 49
 Leu Leu Ile Arg Asn Ile Phe Leu Lys
 1 5
 10
 <210> 50
 <211> 9
 <212> PRT
 <213> Artificial Sequence
 15
 <400> 50
 Gly Leu Gln Glu Phe Asp Ile Ala Arg
 1 5
 20
 <210> 51
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 <400> 51
 25 Lys Met Phe Leu Ile Glu Lys Leu Ile Lys
 1 5 10
 <210> 52
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 30
 <400> 52
 35 Ile Met Leu Phe Ile Met Phe Tyr Ser Lys
 1 5 10
 <210> 53
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 40
 <400> 53
 Lys Leu Ile Lys Leu Gln Asp Met Glu Lys
 1 5 10
 45
 <210> 54
 <211> 10
 <212> PRT
 <213> Artificial Sequence
 50
 <400> 54
 Val Leu Leu Ile Arg Asn Ile Phe Leu Lys
 1 5 10
 55
 <210> 55
 <211> 9
 <212> PRT
 <213> Artificial Sequence
 60
 <400> 55
 Met Pro Arg His Glu Val Tyr Val Leu
 1 5
 65
 <210> 56
 <211> 10
 <212> PRT

```

      <213> Artificial Sequence

      <400> 56
5      Met Pro Arg His Glu Val Tyr Val Leu Leu
      1              5              10

      <210> 57
      <211> 9
      <212> PRT
10     <213> Artificial Sequence

      <400> 57
      Tyr Glu Met Pro Arg His Glu Val Tyr
      1              5

15     <210> 58
      <211> 10
      <212> PRT
      <213> Artificial Sequence

20     <400> 58
      Trp Glu Thr Leu Ile Gly Gln Asp Ile Tyr
      1              5              10

25     <210> 59
      <211> 9
      <212> PRT
      <213> Artificial Sequence

30     <400> 59
      Val His Phe Phe Phe Ile Leu Thr Leu
      1              5

35     <210> 60
      <211> 9
      <212> PRT
      <213> Artificial Sequence

40     <400> 60
      Leu Arg Arg Ile Ile Gly Met Gln Leu
      1              5

45     <210> 61
      <211> 9
      <212> PRT
      <213> Artificial Sequence

      <400> 61
50     Leu Leu Met Asp Phe Val Phe Ser Leu
      1              5

      <210> 62
      <211> 9
      <212> PRT
55     <213> Artificial Sequence

      <400> 62
      Met Gln Leu Ile Thr Ser Leu Gly Leu
      1              5

60     <210> 63
      <211> 9
      <212> PRT
      <213> Artificial Sequence

65     <400> 63

```

```

Phe Val Phe Ser Leu Val Asn Ser Phe
1          5

    <210> 64
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 64
10 Phe Ile Leu Thr Leu Ile Val Leu Ile
   1          5

    <210> 65
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 65
20 Trp Gln Ile Thr Glu Gly Arg Lys Ile
   1          5

    <210> 66
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 66
30 Met Phe Tyr Ser Lys Asn Ile Ser Leu
   1          5

    <210> 67
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 67
35 Phe Ile Met Phe Tyr Ser Lys Asn Ile
   1          5

    <210> 68
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 68
45 Trp Arg Ala Ser Gln Met Met Thr Phe
   1          5

    <210> 69
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 69
55 Phe Phe Phe Ile Leu Thr Leu Ile Val
   1          5

    <210> 70
    <211> 9
    <212> PRT
    <213> Artificial Sequence

    <400> 70
65 Phe Arg Leu Val Glu Arg Tyr Glu Met
   1          5

```

5 <210> 71
 <211> 9
 <212> PRT
 <213> Artificial Sequence

 <400> 71
Phe Phe Ile Leu Thr Leu Ile Val Leu
 1 5

10 <210> 72
 <211> 9
 <212> PRT
 <213> Artificial Sequence

15 <400> 72
Tyr Ala Gln Thr Leu Val Trp Ile Ala
 1 5